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Attorney Docket: 19400/09003 (MJ729)

TECH CENTER IBNIMA USE OF [DOCOSAHEXANOIC] DOCOSAHEXAENOIC ACID AND ARACHIDONIC ACID ENHANCING THE GROWTH OF PRETERM **INFANTS**

FIELD OF INVENTION

The present invention concerns enhancing the growth of preterm infants involving administration of infant formula containing a combination of docosahexaenoic and arachidonic acid.

BACKGROUND OF THE INVENTION

The long chain polyunsaturated fatty acids (LC PUFA) have been shown to be important in infant development. Particularly, arachidonic acid (ARA) and docosahexaenoic acid (DHA) are LC PUFA that are of special interest in infant nutrition because they are found in high concentrations in the brain (Sastry PS, Lipids of nervous tissue: composition and metabolism. Progress Lipid Res 1985;24:69-176) and the retina (Fliesler SJ and Anderson RE. Chemistry and metabolism of lipids in the vertebrate retina. Progress Lipid Res 1983;22:79-131). ARA (20:4n-6) and DHA (22:6n-3) are derived from the parent essential fatty acids linoleic acid (18:2n-6) and α-linolenic acid (18:3n-3) through alternate desaturation and elongation and accumulate rapidly in fetal neural tissue during the last months of gestation and the first months of postnatal life (Makrides M, Neuman MA, Byard RW, Simmer K, Gibson RA. Fatty composition of the brain, retina and erythrocytes in breast- and formula-fed infants. Am J Clin Nutr 1994;60:189-94).

Unlike term infants, preterm infants do not fully benefit from the maternal and placental LC PUFA supply during the last trimester of pregnancy. Even though preterm infants are capable of synthesizing both DHA and ARA from their 18 carbon precursors (Carnielli VP, Wattimena DJL, Luijendijk IHT, Boerlage A, Degenhart HJ, Sauer PJJ. The very low birth weight premature infant is capable of synthesizing arachidonic and docosahexaenoic acids from linoleic and linolenic acids. Pediat Res

1996;40:169-174), it remains unclear whether the rate of synthesis is adequate to meet the optimal needs for central nervous system accretion in the absence of a dietary supply of these fatty acids. Preterm infants are dependent on their own dietary supply of linoleic and α-linolenic acids through either human milk, which also contains small but significant amounts of ARA and DHA or through commercially available artificial formulas, none of which in the United States contain ARA end DHA.

It has been demonstrated in recent studies (Hoffman DR and Uauy R. Essentiality of dietary ω -3 fatty acids for premature infants: Plasma and red blood cell fatty acid composition. Lipids 1992;27:886-95) that the fatty acid composition of red blood cell membrane lipids in infants receiving formulas supplemented with DHA (0.35% of total fatty acids) was similar to human milk-fed infants. In the same study, Birch (Birch DG, Birch EE, Hoffman DR Uauy RD. Retinal development in very-low-birth-weight infants fed diets differing in Omega-3 fatty acids. Investigation Ophthalmology Visual Science 1992;33:2365-76) found that retinal function improved with the provision of a dietary supply of DHA in very low birth weight infants.

The first year growth of preterm infants fed standard formula compared to marine oil LC PUFA supplemented formula was studied by Carlson et al. (Carlson SE, Cooke, RJ, Werkman SH, Tolley EA. First year growth of preterm infants fed standard compared to marine oil n-3 supplemented formula Lipids 1992:27:901-907). The experimental formulas provided 0.2% of total fatty acids as DHA and also provided 0.3% as EPA (20:5n-3). This EPA concentration is higher than found in human milk while the DHA level is similar to human milk. Beginning at 40 weeks from conception, marine oil supplemented infants compared to controls had significantly lower weight, length, and head circumference. From this study, Carlson (Carlson SE, Werkman SH, Peeles JM, Cooke RJ, Tolley EA. Arachidonic acid status correlates with first year growth in

preterm infants. Proc Natl Acad Sci USA 1993;90:1073-77) hypothesized that dietary ARA could improve first year growth of preterm infants, in the context of restoring growth to the level of control formula containing no LC PUFA.

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In another study (Montalto, FB, et al., Pediatric Research, Vol 39, page 316A, abstract no. 1878) it was shown that male infants fed marine oil supplemented formula (containing DHA but essentially no ARA) had, by 4 to 6 months, lower head circumference, length, weight and fat free mass than standard formula fed infants. A third study also showed decreased weight at 9 and 12 months corrected age in preterm infants fed marine oil supplemented formula (with LC PUFA) to 2 months corrected age compared with control formula containing no LC PUFA (Carlson SE, et al., Am. J. Clin. Nutr., 63 pp 687-97, 1996).

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The prior art has demonstrated that infants with altered tissue LC PUFA levels, resulting from a lack of LC PUFA in their diets, may be at risk for neurological problems, may also have reduced scores on cognitive tests, and may have lower retinal development than human milk-fed infants. Worldwide regulatory organizations such as the WHO/FAO Expert Committee on Fats and Oils in Human Nutrition have recommended that LC PUFA be included in preterm infant formula. These recommendations have been made despite the negative effects observed of DHA supplements on growth. There has been no demonstration in the literature that ARA and DHA, particularly when added to infant formula, enhances the growth of infants above that demonstrated by control formulas not containing ARA and DHA.

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SUMMARY OF THE INVENTION

It has unexpectedly been discovered that preterm infants receiving infant formula supplemented with both DHA and ARA demonstrate enhanced growth. The present invention is directed to enhancing the growth of preterm infants comprising administering to said infants a growth

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enhancing amount of DHA and ARA.

DETAILED DESCRIPTION OF THE INVENTION

As reported in a review of preterm infant growth by Carlson, SE, (The Jrnl of Pediatrics, vol 125, pp 533-8, 1994) "After adjusting for postconceptional age, preterm infants show a decline (rather than a catch-up) in the normalized weight from approximately 2 to 4 months past expected term."

Several prior art studies have documented the value of administering DHA to infants. However, when DHA, either as the primary LC PUFA or combined with EPA, is administered to preterm infants, said infants suffer from decreased growth. It has been suggested that ARA may be beneficial to growth; however, heretofore the growth effects of administering both DHA and ARA to preterm infants have been unknown. It has been surprisingly discovered that administering the combination of ARA and DHA results in enhanced growth of infants relative to infants fed DHA alone. It has also been discovered that preterm infants administered an infant formula containing ARA and DHA exhibit enhanced growth relative to preterm infants fed control formula without DHA and ARA, such as those formulas currently used in modern nurseries. It has further been discovered that practice of the method of the invention results in growth of preterm infants catching up in an unexpected short time to a reference group of normal term breast fed infants.

The time to achieve growth similar or equivalent to normal term breast fed infants by practice of the method of the invention is less than 9 months corrected age; preferably less than 6 months corrected age, more preferably less than 4 months corrected age, even more preferably less than 2 months corrected age, and most preferably no greater than term corrected age.

The method of the invention requires a combination of DHA and ARA. The weight ratio weight of ARA:DHA can be about 1:2 to about 5:1,

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preferably about 1:1 to about 3:1, and more preferably about 2:1.

In the method of the invention the combination of DHA and ARA is preferably administered as part of an infant formula. The infant formula for use in the present invention is preferably nutritionally complete and typically contains suitable types and amounts of lipid, carbohydrate, protein, vitamins and minerals. The amount of lipid or fat typically can vary from about 3 to about 7 g/100 kcal. The amount of protein typically can vary from about 1 to about 5 g/100 kcal. The amount of carbohydrate typically can vary from about 8 to about 12 g/100 kcal. Protein sources can be any used in the art, e.g., nonfat milk, whey protein, casein, soy protein, hydrolyzed protein, amino acids, and the like. Carbohydrate sources can be any used in the art, e.g., lactose, glucose, corn syrup solids, maltodextrins, sucrose, starch, rice syrup solids, and the like. Lipid sources can be any used in the art, e.g., vegetable oils such as palm oil, soybean oil, palmolein, coconut oil, medium chain triglyceride oil, high oleic sunflower oil, high oleic safflower oil, and the like. Conveniently, commercially available infant formula can be used. For example, Enfamil®, Enfamil® Premature Formula, Enfamil® with Iron, Lactofree®, Nutramigen®, Pregestimil®, ProSobee® (available from Mead Johnson & Company, Evansville, Indiana, U.S.A.), Similac®, Isomil®, Alimentum®, Neocare®, and Similac® Special Care (available from Ross Laboratories, Columbus, Ohio, U.S.A.), may be supplemented with suitable levels of ARA and DHA at the proper ratios and used in practice of the method of the invention.

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The form of administration of the DHA and ARA in the method of the invention is not critical, as long as a growth enhancing amount is administered. Most conveniently, the DHA and ARA are supplemented into infant formula which is then fed to the infants. Alternatively, the DHA and ARA can be administered as a supplement not integral to the formula feeding, for example, as oil drops, sachets, in combination with other

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nutrient supplements such as vitamins, and the like.

The growth enhancing amount of DHA is typically about 2.5 mg/kg of body weight/day to about 60 mg/kg of body weight/day, preferably about 6 mg/kg of body weight/day to about 40 mg/kg of body weight/day, more preferably about 12 mg/kg body weight/day to about 30 mg/kg body weight/day, and even more preferably about 18 mg/kg of body weight/day to about 24 mg/kg of body weight/day.

The growth enhancing amount of ARA is typically about 5 mg/kg of body weight/day to about 120 mg/kg of body weight/day, preferably about 12 mg/kg of body weight/day to about 80 mg/kg of body weight/day, more preferably about 24 mg/kg body weight/day to about 60 mg/kg body weight/day, and even more preferably about 36 mg/kg of body weight/day to about 48 mg/kg body weight/day.

The amount of DHA in infant formulas for use in the present invention typically varies from about 2 mg/100 kilocalories (kcal) to about 50 mg/100 kcal, preferably about 5 mg/100 kcal to about 33 mg/100 kcal, more preferably about 10 mg/100 kcal to about 25 mg/100 kcal, and even more preferably about 15 mg/100 kcal to about 20 mg/100 kcal.

The amount of ARA in infant formula for use in the present invention typically varies from about 4 mg/100 kcal to about 100 mg/100 kcal, preferably about 10 mg/100 kcal to about 67 mg/100 kcal, more preferably about 20 mg/100 kcal to about 50 mg/100 kcal, and even more preferably about 30 mg/100 kcal to about 40 mg/100 kcal.

The infant formula supplemented with oils containing DHA and ARA for use in the present invention can be made using standard techniques known in the art. For example, replacing an equivalent amount of an oil normally present, e. g., high oleic sunflower oil.

The source of the ARA and DHA can be any source known in the art such as fish oil, single cell oil, egg yolk lipid, brain lipid, and the like.

The DHA and ARA can be in natural form, provided that the remainder of

the LC PUFA source does not result in any substantial deleterious effect on the infant. Alternatively, the DHA and ARA can be used in refined form. It is preferred that the LC PUFA used in the invention contain little or no EPA. For example, it is preferred that the infant formulas used herein contain less than about [10 mg/100 kcal] 20 mg/100 kcal EPA; preferably less than about 10 mg/100 kcal EPA; more preferably less than about 5 mg/100 kcal EPA; and most preferably substantially no EPA.

Preferred sources of DHA and ARA are single cell oils as taught in U.S. patent nos. 5,374,657, 5,550,156, and 5,397,591, the disclosures of which are incorporated herein by reference in their entirety.

The following examples are to illustrate the invention but should not be interpreted as a limitation thereon.

EXAMPLES

I CLINICAL STUDY DESIGN

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1. INTRODUCTION

This study is a double-blind, randomized, controlled parallel design, prospective trial of premature infant formulas containing microalgae and fungi-derived oils which contain a part of their constituents arachidonic acid and docosahexaenoic acid. Formula feeding subjects will be randomized into one of 3 feeding groups:

- premature formula plus DHA (about 0.13% of energy)
 and ARA (about 0.26% of energy)
- premature formula plus DHA (about 0.13% of energy)
- premature formula WITHOUT DHA and ARA

The products have the same nutrient composition (see Appendix A) and differ only in the level of DHA and ARA. The products will be blinded. The present order of formula has no relationship to randomization.

Normal, term, breast fed infants will be enrolled to provide a normal visual acuity reference.

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Fifty evaluable subjects will be completed in each group. Premature infants will remain on study formulas after reaching 90 kcal/kg/d for a minimum of 28 days or until hospital discharge whichever is longer. After 28 days or discharge, whichever is longer, all premature infants will receive Enfamil or Enfalac with Iron. If medically indicated, ProSobee, Lactofree, Alactamil, Nutramigen, or Pregestimil may be used in place of Enfamil or Enfalac with Iron. Term infants will receive at least 85% of their nutrition from breast milk. Primary measures of effectiveness will include visual acuity and red blood cell membrane fatty acid profiles (i.e. DHA and ARA levels). The measure of safety will be growth and adverse experience reports.

2. SUBJECTS

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2.1 SOURCE AND CHARACTERIZATION OF STUDY GROUP

Acceptable preterm subjects will be relatively healthy premature infants taking preterm formula. Anticipated hospitalization should be sufficient to allow for 28 days of enteral intake \geq 90 kcal/kg/d and \geq 85% study formula intake. All races and both sexes will be eligible for the study.

2.2. INCLUSION CRITERIA

20 Preterm infants:

- Birth weight ≥ 900 g
- Formula feeding at time of study enrollment
- Anticipate enteral intake of ≥90 kcal/kg/day for ≥ 28 days before discharge home
- Informed consent obtained

Term Infants:

- 38 to 42 weeks gestation
- Committed to breast feeding
- Informed Consent obtained

30 2.3 EXCLUSION CRITERIA

Preterm infants:

• ≥ 1500 g at birth

Preterm and Term Infants:

- History of underlying disease or congenital malformation which in the opinion of the investigator is likely to interfere with the evaluation of the subject
- More than 24 days between birth and full oral feeds (≥ 90 kcal/kg/d)
- Small (<10th percentile) for gestational age at birth (SGA)
- Necrotizing enterocolitis as diagnosed by the physician
- Other gastrointestinal disease
- Impaired visual or ocular status at birth

2.4 CONCOMITANT MEDICATIONS, HOSPITALIZATIONS,

ILLNESSES

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- No medication which may affect FPL response may be used within 3 days of measurement.
- No evidence of viral of bacterial infection during FPL testing.
- No medications known to [effect] <u>affect</u> lipid metabolism (e.g., heparin at therapeutic levels)

20 3. STUDY PRODUCT INFORMATION

3.1 FORMULATIONS

Nutrient composition is included as Appendix A.

4. STUDY PROCEDURES

4.2.1 ENROLLMENT

25 Enrollment will take place over a 6 month period. Ideally, sufficient subjects will be enrolled so that 10 subjects in each group complete the study at each site for the multi-center trial. A total of 50 infants per formula group will complete this trial.

4.2.2 SCHEDULE OF EVENTS (SEE FLOW CHART, SECTION

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4.2.2.1 RECRUITMENT

Mothers of eligible, healthy, preterm formula fed infants and term, breastfed infants will be contacted, the study explained to them, and if they are agreeable, written informed consent obtained.

Term infants may be enrolled anytime from birth until or during the 48 week visit.

4.2.2.2 RANDOMIZATION

Recruited formula fed subjects will be randomized into study groups. Randomization can occur anytime after enteral feeds reach 50 kcal/kg/day until commencement of full enteral feeds (i.e., ≥90 kcal/kg/day).

4.2.2.3 **FEEDING**

All premature infants will receive their assigned study formula after informed consent has been granted and enteral feeds are at least 50 kcal/kg/day. The infant will remain on study formula 28 days after reaching 90 kcal/kg/d or until hospital discharge, whichever is longer. Oral feeding amount, strength and rate will advance as appropriate for the clinical management of the infant.

All parents will be instructed not to feed solid foods during the study. The parents will be instructed that the study formula or breast milk is to serve as the sole source of food from enrollment to study end.

4.2.2.4 BASELINE DATA COLLECTION

The following data will be collected by the Investigator at the time of enrollment and randomization on the case report forms:

- Informed consent of parent obtained.
- Post conceptual age.
- That the subject is a premature infant, with Birth weight
 ≥900 gm and ≥1500 gm or a normal term infant between
 38 and 42 weeks gestational age.
 - That the preterm subject is receiving infant formula or

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term infant is committed to breast feeding. Anticipated preterm infant enteral intake of ≥90 kcal/kg/day for ≥28 days prior to discharge home. That the subject has no history of underlying disease, inborn error of metabolism, or congenital malformation 5 which in the opinion of the Investigator is likely to interfere with the evaluation of the study formulas. That the subject is not small (<10th percentile) for gestational age at birth. 10 That the subject does not have necrotizing enterocolitis as diagnosed by a physician. That the subject does not have a gastrointestinal disease. No more than 24 days between birth and full enteral 15 feeds (i.e., ≥90 kcal/kg/day). That the subject did not have impaired visual or ocular status at birth. Birth date, sex, race. Birth weight, length and head circumference 20 4.2.2.5 INVESTIGATOR PERIODIC DATA COLLECTION "During hospitalization, preterm subjects will have their weight recorded daily while they are receiving study formula. Length and head circumference will be recorded weekly, along with an additional weight measurement. For a given subject, the same scale should be used for the weekly 25 weight measurement."

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4.2.2.6 BLOOD DRAW

"Weight, length, and head circumference will also be

recorded at the 40, 48, and 57 week post conceptual age

visit (preterm) and 56 and 119 days of age visit (term)."

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When preterm infant enrolls in the study and again at termination of study formula (i.e., hospital discharge or 28 days after reaching 90 kcal/kg/d of study product), the Investigator will ascertain that the infant is essentially solely formula fed. If this criteria is met, 1.2 ml/blood will be drawn for blood lipids. The sample will be processed as described in Appendix B.

An attempt will also be made to draw a similar blood sample at the 48 weeks PCA visit when visual acuity is measured in both term and preterm infants.

4.2.2.7 VISUAL ACUITY BY FORCED CHOICE PREFERENTIAL LOOKING (FPL) AT 48 AND 57 WEEKS \pm 4 DAYS POST-CONCEPTUAL AGE

When the infant is 48 and 57 weeks \pm 4 days post-conceptual age, trained persons at each study site will follow the Teller Acuity Card Procedure for the measurement of visual acuity of all study subjects. It is essential that only persons who are trained in the FPL procedure for determining visual acuity do the testing. If necessary, training of responsible persons and documentation of completion of successful training will be done at Children's Hospital Medical Center Ophthalmology Department in Seattle, Washington, according to the procedure attached as Appendix C.

If the infant cannot complete the procedure at 48 or 57 weeks \pm 4 days postconceptual age (i.e., too fussy, too sleepy, too inattentive) the test should be repeated within 7 days.

4.2.2.8 INTERIM EVALUATION

At preterm infant hospital discharge or 28 days after reaching 90 kcal/kg/d of study formula feeding, whichever is longer, the [investigator] Investigator will fill out an "Interim Evaluation" form. After reviewing the subject's records and discussion with the parents and staff, the [investigator will indicate whether:] Investigator will indicate:

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- Whether or not the subject completed at least 28 days of study formula intake ≥ 90 kcal/kg/d and both blood samples obtained
- If the study was not completed, and reason
- Whether or not the subject received steroids (glucorticoids)
- Investigator's evaluation of the study formula

The first and last dates study material was taken will be recorded.

4.2.2.9 FINAL EVALUATION

At the final study visit (57 weeks postconceptual age) or earlier if the subject drops out, the Investigator will fill out a "Final Evaluation" Case Report Form. After reviewing the subject's records and discussion with the parents, the Investigator will indicate whether the subject:

- (1) Completed feeding regiment and all study parameters (i.e., anthropometrics and visual acuity measured).
 - (2) Did not complete feeding regimen.
 - (3) Not completed and reason.

4.3 CLINICAL OBSERVATIONS

4.3.1 PHYSICAL EXAMINATIONS

Subjects will have weight, length and head circumferences recorded at birth, weekly while hospitalized, then at 40, 48, and 57 weeks \pm 4 days postconceptual age.

Body weight will be measured using an electronic balance or a double beam balance accurate to 10 g or ½ oz with non-detachable weights. During hospitalization, if more than one such balance is employed in the practice, either one balance should be designated the study balance and all study weights will be carried out on that balance for a particular subject, or the balances will be checked and certified to register the same weight throughout the range of weights expected.

Outpatient weights will be obtained on a calibrated office scale.

Documentation indicating balance calibration of the outpatient balance carried out within 12 months of study initiation will be supplied to the Sponsor.

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Length will be measured with the infant in recumbent position with the help of two examiners and a suitable measuring apparatus. One person holds the subject's head in contact with a fixed vertical headboard and a second person holds the subject's feet, toes pointing directly upward and, also applying gentle traction. The baby is measured from the headboard to the soles of the feet with a non-stretching tape measure.

Head circumference will be measured, employing a flexible, non-stretchable cloth or vinyl tape.

4.3.2 VISUAL ACUITY BY FORCED CHOICE PREFERENTIAL 15 LOOKING (FPL)

Visual acuity will be determined at 48 and 57 weeks \pm 4 days postconceptual age according to procedures outlined in Appendix C.

4.3.3 LABORATORY TESTS

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Blood will be drawn from preterm infants by heel prick or venipuncture when study formula is begun and terminated. An attempt will be made to draw blood at 48 weeks \pm 4 days PCA from both term and preterm infants. Procedures for handling the blood are described in Appendix B.

4.4 FLOW CHART

			PREI	PRETERM				TERM	
EVENT	# BIMP	Enteral Intake >50 kcal/kg/d	Termination of Study Formula t	Visit 1 40 wks ± 4d PCA	VISIT 2 46 wks ± 4d PCA	Visit 3 57 wks ± 4d PCA	Visit 1 40 wks ± 44 PCA	Visit 2 46 wks ± 4d PCA	Visit 3 57 wke ± 44 PCA
Randomization		>							
Study Formula		>							
Enfamil w/iron	nene.	·	>	>	>	>			
Human Milk	Ţ						>	>	>
	<u>.</u>		Phy	Physical		·		Physical	
Weight	` >	*	>	>	>	>	>	>.	>
Lèngth	· >	*>	>	>	>	>	<i>></i>	>	>
Head Circumference	>	\$	>	>	>	>	>	>	>
Blood Draw		>	>	·	^			>	
Visual Acuity Test	. •- = ≥			·	>	>		>	>
Illnesses	N	·		>	>	>		>	>
Interim Assessment			>				· ·		
Final Assessment	The second second second)	(when the subject discontinues or completes)	continues or comp	oletes}		(when the sub	(when the subject discontinues or completes)	completes)

Medical problems related to or affecting formula consumption will be recorded when they occur.
 Recorded daily and weekly during hospitalization.
 At hospital discharge or 28 days of study formula intake (after reaching 90 kcal/kg/d), whichever is later.

5. CRITERIA FOR RESPONSE

Criteria for response will depend upon the following:

- Visual Acuity better than the control formula.
- Visual Acuity comparable to breastfed term infant.
- Red Blood Cell phosphatidyl ethanolamine DHA and ARA weight % greater than formula control group.
- Growth as measured by weight achieved at 48 and 57 weeks postconceptual age comparable to formula control group.

10 **6. STATISTICS**

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6.1 RANDOMIZATION

If the subject meets the inclusion and exclusion criteria, randomization to one of three formula groups will take place. The randomization schedule will be provided by Mead Johnson Research Center. A separate randomization schedule will be provided for males and females.

6.2 SAMPLE SIZE

The primary parameter of interest is visual acuity as measured by the Forced Choice Preferential Looking (FPL). The minimal clinically relevant difference was determined to be 0.5 octave. A consultant in the field of visual acuity estimated the standard deviation to be 0.5 octave. This value was increased to .7 octave in case more variability was experienced in this study. Thirty-two subjects per group are needed to attain 80% power when testing at an alpha level of 0.05.

A sample size estimate of 50 per group was determined to achieve α + 0.05, β + 0.20, for weight of infants receiving study oil being greater than 400 gm below control at 48 weeks postconceptual age or 500 g below control at 57 weeks postconceptual age with a standard deviation of 800 g. It was therefore determined that 50 subjects per group will be used in the study.

6.3 ANALYTICAL PLAN

Visual acuity data will be recorded in cycles per cm. These values will be converted to cycles per degree using the following formula:

cycles/degree =
$$\frac{38 \text{ x cycles/cm}}{55}$$

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A log transformation will be applied to the data prior to analysis. Analysis of variance techniques will be used to assess feeding regimen group differences in visual acuity. If the overall F test for feeding regimen is significant at [al] an alpha level of 0.05, pairwise comparisons will be made at an alpha level of 0.05. If no significant differences are detected, then a post-study power analysis will be performed to demonstrate that the study had adequate power to detect the minimal clinically relevant difference.

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Analysis of variance will be used to assess feeding regimen differences in phosphatidyl choline DHA and ARA levels and in phosphatidyl ethanolamine DHA and ARA levels at each time point. If the overall F test is significant at [al] <u>an</u> alpha level of 0.05, then pairwise comparisons will be made at an alpha level of 0.05.

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Analysis of variance will be used to assess feeding regiment differences in weight at 48 and 57 weeks postconceptual age. The statistical model will include terms for feeding regimen, study center, sex and all two-way interactions. Non-significant interactions will be removed from the final statistical model. Two one-sided tests will be performed comparing each experimental formula (EC) with the control formula (CF). The hypothesis to be tested is as follows:

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$$H_0$$
 = Weight (CF) \leq Weight (EF).

The alternative hypothesis is as follows:

$$-H_1$$
 = Weight (CF) > Weight (EF).

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If H_0 [if rejected] <u>is rejected</u> and the mean weight of the control formula exceeds that of the experimental formula by more than

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400 mg at 48 weeks postconceptual age or by 500 g at 57 weeks postconceptual age then the conclusion is that the experimental formula does not exceed that of the experimental formula by more than 400 g at 48 weeks postconceptual age or by 500 mg at 57 weeks postconceptual age then the conclusion is that the experimental formula does provide adequate growth. If H₀ is not rejected then a post-study power analysis will be performed to demonstrate that [eh] the study had adequate power to detect the above mentioned clinically relevant differences. If adequate power is achieved then the conclusion is that the experimental formula does provide adequate growth.

Fisher's exact test will be used to compare the proportion of subjects in each group with illness/symptoms of concern during the study. The analysis will be performed for each type of illness/symptom reported, with classification of investigator terms into similar terminology made as necessary.

APPENDIX A NUTRIENT COMPOSITION OF FORMULAS

All study formulas are 24 kcal/fl oz and are identical in composition to marketed Enfamil Premature Formula except for the study oils employed. These oils are described in the protocol.

NUTRIENT AMOUNT/100 kcal ENFAMIL WITH Fe Protein g 3 2.2 Fat, g 5.1 5.6 Carbohydrate, g 11.1 10.3 Vitamin A IU 1250 310 Vitamin D IU 270 63 Vitamin E IU 6.3 .2 Vitamin K mcg 8 8 Thiamine, mcg 200 78 Riboflavin, mcg 300 150 Vitamin B ₀ , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5		STUDY FORMULAS	
Fat, g 5.1 5.6 Carbohydrate, g 11.1 10.3 Vitamin A IU 1250 310 Vitamin D IU 270 63 Vitamin E IU 6.3 .2 Vitamin K mcg 8 8 Thiamine, mcg 200 78 Riboflavin, mcg 300 150 Vitamin B ₆ , mcg 150 63 Vitamin B ₁₂ mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 Magnesium, mg 6.3 Iron, mg 1.8 Vitamin C, mg 6.3 Magnesium, mg 1.8 Iron, mg 1.8	NUTRIENT	AMOUNT/100 kcal	ENFAMIL WITH Fe
Carbohydrate, g 11.1 10.3 Vitamin A IU 1250 310 Vitamin D IU 270 63 Vitamin E IU 6.3 .2 Vitamin K mcg 8 8 8 Thiamine, mcg 200 78 Riboflavin, mcg 150 63 Vitamin B ₆ , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 Magnesium, mg 6.3 Iron, mg 1.8	Protein g	3	2.2
Vitamin A IU 1250 310 Vitamin D IU 270 63 Vitamin E IU 6.3 .2 Vitamin K mcg 8 8 Thiamine, mcg 200 78 Riboflavin, mcg 300 150 Vitamin B ₆ , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, meg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Fat, g	5.1	5.6
Vitamin D IU 270 63 Vitamin E IU 6.3 .2 Vitamin K mcg 8 8 Thiamine, mcg 200 78 Riboflavin, mcg 300 150 Vitamin B ₆ , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Carbohydrate, g	11.1	10.3
Vitamin E IU 6.3 .2 Vitamin K mcg 8 8 Thiamine, mcg 200 78 Riboflavin, mcg 300 150 Vitamin B ₆ , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Vitamin A IU	1250	310
Vitamin K mcg 8 8 Thiamine, mcg 200 78 Riboflavin, mcg 300 150 Vitamin B ₆ , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Vitamin D IU	270	63
Thiamine, mcg 200 78 Riboflavin, mcg 300 150 Vitamin B ₆ , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 6.3 7.8 Iron, mg 1.8 0.5	Vitamin E IU	6.3	.2
Riboflavin, mcg 300 150 Vitamin B _s , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Vitamin K mcg	8.	8
Vitamin B ₆ , mcg 150 63 Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Thiamine, mcg	200	78
Vitamin B ₁₂ , mcg 0.25 0.23 Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Riboflavin, mcg	300	150
Niacin, mcg 4000 1250 Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Vitamin B ₆ , mcg	150	63
Folic Acid, mcg 35 15.6 Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Vitamin B _{12,} mcg	0.25	0.23
Pantothenate, mcg 1200 470 Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Niacin, mcg	4000	1250
Biotin, mcg 4 2.3 Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Folic Acid, mcg	35	15.6
Vitamin C, mg 20 8.1 Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Pantothenate, mcg	1200	470
Choline, mg 12 15.6 Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Biotin, mcg	4	2.3
Inositol, mg 17 4.7 Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Vitamin C, mg	20	8.1
Calcium, mg 165 78 Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Choline, mg	12	15.6
Phosphorus, mg 83 53 Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Inositol, mg	17	4.7
Magnesium, mg 6.3 7.8 Iron, mg 1.8 0.5	Calcium, mg	165	78
Iron, mg 1.8 0.5	Phosphorus, mg	83	53
, 6	Magnesium, mg	6.3	7.8
Zinc, mg 1.5 0.78	Iron, mg	1.8	0.5
	Zinc, mg	1.5	0.78

	STUDY FORMULAS	
NUTRIENT	AMOUNT/100 kcal	ENFAMIL WITH Fe
Manganese, mcg	6.3	15.6
Copper, mcg	125	94
lodine, mcg	25	6
Sodium mg (mEq)	39 (1.7)	27 (1.17)
Potassium mg(Meq)	103 (2.6)	108 (2.8)
Chloride mg (Meq)	85 (2.4)	63 (1.77)

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FINAL STUDY REPORT

Study Design:

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This double-blind, parallel-group study (project 3338) was carried out in 16 neonatal centers (study numbers 9698-9709, 9712, 9723, 9743, and 9746) in North America. Three premature infant feedings were compared. Each had the same composition except for the incorporation of fungal and/or micro algal oils up to about 3% of the fat blend to provide the experimental levels of docosahexaenoic acid (DHA) and arachidonic acid (ARA). The control formula (C, Enfamil® Premature Formula) contained no DHA or ARA, the DHA formula (D) contained about 0.15% of energy as DHA (0.34% of fat), and the DHA+ARA formula (DA) contained about 0.14% of energy as DHA (0.33% of fat) and 0.27% of energy as ARA (0.60% of fat). The formulas were fed to 284 randomized infants weighing 846 to 1560 grams at birth for at least 28 days. Upon completion of study formula intake, they were given routine infant formula and followed through 4 months gestationally corrected age. A group of 90 exclusively human milk fed term infants were enrolled and followed to 4 months of age as a reference group (H).

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Study Objective and Statistical Analysis:

The primary objective of this study was to establish the safety of

feeding D or DA to preterm infants during their initial hospitalization as measured 1) by growth, acceptance and tolerance while consuming the formula for at least 1 month and 2) by close monitoring and observation for a 4 to 5 month follow-up period (4-5 times the treatment period) while consuming unsupplemented routine term infant formula. The primary growth parameter selected was weight with evaluation of the proposition that weight on test formula was greater than or equal to weight on control formula. The one sided statistical test for an adverse effect on growth maximized the power to detect a difference should one be present. A two-sided test was used for all other parameters. A p-value of less than 0.05 was used to establish significance.

Secondary objectives of the study were 1) to evaluate the impact of fatty acid levels in erythrocyte phospholipids at the end of study feeding and 2) to determine if any effect on mean visual acuity greater than half an octave could be demonstrated at 2 and 4 months corrected age.

Results:

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Six infants were just outside the weight parameters and five infants just older than the less than 24 days chronological age parameter for enrollment in the study. In each case, judgement by the clinical or medical monitor was made to include them in the study prior to enrollment based on their homogeneity with other study infants in all other particulars, e.g., state of health, type of medical complications, and weight for gestational age. All these infants were included in the analysis of the study results.

The formula groups were comparable at enrollment (See table 1). Post-conceptual age, weight, length, and head circumference at enrollment did not differ among the groups.

All groups experienced comparable final study status (See table 2).

Drop outs did not differ among the formula fed groups during hospitalization. There also were no differences in drop outs among the four groups at study completion.

Both formulas D and DA provide adequate growth when compared to formula C (See table 3, figure 1, and Appendix 1). Weight gain during hospitalization was no less on D or DA than on C, 33.3, 34.7, and 30.7 g/day, respectively. Furthermore, no less weight was achieved on D or DA than on C at 40, 48, and 57 weeks post-conceptual age (See table 4, figure 2, and Appendix 1); statistical power was greater than 0.89 to detect a clinically relevant decrease.

Post-hoc analysis reveals that infants on DA grew faster than infants receiving C and D (See table 5 and figure 1). This enhanced growth provided faster "premature infant catch-up" compared to C and D. Weight achieved by the DA group (3198 g) was higher than C (3075 g) and D (3051 g) at 40 weeks post-conceptual age but had not fully caught up to the term birth weight (3438 g) of group H (See table 4 and figure 2). This catch up trend continued through 48 to 57 weeks by which time the mean weight of group DA did not differ from group H while groups C and D remained significantly lower.

Length was not different among the formula groups either during hospitalization or the follow-up period, although the ordered sequence of mean lengths was the same as for the weights (See table 7 and figure 3). This is likely at least partially due to length being a less sensitive parameter of growth than weight. For the same reason, the mean lengths of group H infants were higher than that of all the premature infant groups at 40, 48 and 57 weeks post-conceptual age indicating slower catch up in this parameter.

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Head circumference is the least sensitive parameter of growth and was not different among any of the four groups at any time measured except at 40 weeks postconceptual age (See table 8 and figure 4). At this time, as expected, the birth head circumference of group H was smaller than the formula fed premature infants possibly due to molding of labor and to insufficient time for adjustment to the extrauterine environment.

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Visual acuity has reportedly been enhanced in studies where DHA supplemented formulas were fed to premature infants both in the hospital and continuing after discharge. In this study, visual acuity was measured about 3 months and then about 5 months after stopping study formula to determine whether a residual beneficial effect of at least half an octave might be observed. Although no difference in visual acuity was found among the formula groups at these times (See table 8 and figure 5), the acuity card method used, the length of study formula feeding, and/or the length of time not on study formula at the time of measurement may have precluded its detection. However, at 57 weeks post-conceptual age, the breast fed term infant group did have statistically higher visual acuity scores than the test formula groups. But even these differences were at most only 0.33 octave and were clinically insignificant (See figure 6). It is important to note that the breast fed infants continued to receive DHA and ARA during the 3-5 month follow-up period while the formula fed groups did not. Thus, this minor difference in performance was not unexpected based on previous study findings and on developmental differences between term and preterm infants even at the same gestational age.

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Individual fatty acid levels were determined in the phosphatidylcholine and phosphatidylethanolamine fractions of red blood cells before formula feeding, at the conclusion of test formula feeding, and at 48 weeks post-conceptual age (See tables 9 and 10). The premature infant groups were comparable at the beginning of test formula feeding. At the conclusion of test formula feeding, individual fatty acid levels varied among the groups. DHA and ARA were statistically significantly higher in the respectively supplemented groups. Other fatty acid levels reflected the impact of the supplementation. No clinically significant alterations in fatty acid levels or metabolism were identified. After discontinuing study formula and consuming a diet without DHA or ARA for about 3 months, no differences in fatty acid levels among formula fed groups were detectable,

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except for [phosphatidylethanolmine] phosphatidylethanolamine levels of 18:2 (range 8.9-9.3%) and DHA (range 3.2-4.1%) which differences were not identified as being clinically significant. However, the breast fed group shows statistically significant differences in 13 fatty acid levels compared to the formula fed infants. These differences are undoubtedly due to the differences in fatty acid composition of human milk and the term formulas including the lack of DHA and ARA in the latter.

Preterm infant complications were similar in all groups (See table 11). Over 80% of all infants were opthamologically examined and over 90% had ultrasound evaluation of their heads. Specifically, the incidence and severity of retinopathy of prematurity (ROP or retrolental fibroplasia/RLF) and the incidence of intraventricular hemorrhage or its complications did not differ among formula groups. No feeding group related complications were identified.

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Serious adverse experiences did not differ (p = 0.93) among the formula groups and were in the range of those expected in a premature infant population while on study formula: 6% in group C, 5% in group D, and 6% in group DA (See table 12). After the experimental formula phase, serious adverse experiences still did not differ among the preterm groups (See table 13): 13% in group C, 15% in group D, and 15% in group DA. However, the term infant breast fed-group had significantly fewer serious adverse experiences (1%, p = 0.002) as expected. Two infants reportedly suffered sudden infant death syndrome (SIDS), one in group C and one in group D; there was no significant difference in this complication among all four groups.

Conclusions:

We conclude that feeding 0.13% of calories as DHA from micro algal oil and feeding 0.13% of calories as DHA from micro algal oil plus 0.26% of calories as ARA from fungal oil in the matrix of premature infant formula to premature infants during the period of their initial hospitalization

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prior to 40 weeks post conceptual age is safe. These micro algal and fungal oil supplements do not result in any adverse effect on growth, clinical complications, or untoward events. Furthermore, this study reveals that growth benefits accrue to premature infants fed Enfamil Premature Formula supplemented with DHA and ARA from these sources compared to unsupplemented formula or formula supplemented with only DHA. No measurable benefit on visual acuity was identified when infants were tested at about 3 and 5 months after the supplemented formula was discontinued (2 and 4 months corrected age). However, providing human milk levels of intake of long chain polyunsaturated acids are warranted because they are critical to brain development and foster enhanced catchup growth during this early development period.

Table I
Birth Statistics of Premature Subjects

	T n	Mean (std)	D	
Post Conserval Ass (IV. des)	+ "	ivicali (Stu)	Range	p-value
Post-Conceptual Age (Weeks) Control DHA DHA+ARA	62 66 66	29.5 (1.7) 30.0 (1.4) 29.7 (1.7)	25 - 33 26 - 32 26 - 34	0.076
Birth Weight (g) Control DHA DHA+ARA	62 66 66	1233.1 (176.6) 1272.8 (168.1) 1278.9 (177.6)	846 - 1560 900 - 1545 910 - 1535	0.25
Birth Length (cm) Control DHA DHA+ARA	60 66 66	38.4 (2.3) 38.6 (2.2) 38.7 (2.3)	34 - 43.75 33 - 43.5 33 - 44	0.62
Birth Head Circumference (cm) Control DHA DHA+ARA	61 64 65	26.9 (1.5) 27.3 (2.1) 27.2 (1.6)	23.5 - 30.5 22 - 37 23.5 - 30	0.53

Table 2 Summary of Final Study Status

	ļ	R	egimen		
	Control	DHA	DHA+ARA	HM	p-valu
Immediate dropout, study formula never consumed		2	2	HM	
Study Formula Phase *			 	+	
Completed Discontinued	52 (84%) 10 (16%)	59 (89%) 7 (11%)	62 (94%) 4 (6%)		0.20
Reason discontinued					
>96 cumulative hours NPO <28 days of intake >= 90 kcal/kg/day Complications unrelated to study	3 3	1 . 3			
formula NEC or other GI disease Formula intolerance	1	1	1		
Parents request Not off oxygen prior to discharge Protocol violation	2	2	I I	i	-
Term Formula Phase **					
Completed Discontinued	45 (87%)	47 (80%)	53 (85%)	77 (86%)	0.74
Discondinated	7 (13%)	12 (20%)	9 (15%)	13 (14%)	

The CRFs for 9709-003 (DHA) and 9743-304 (DHA) were marked discontinued because the subjects met the study formula intake criteria for only 27 days. These subjects are counted completed here because subjects at other sites with similar intakes were marked completed.

*Based on subjects who completed the Study Formula phase. During the Term Formula phase, subjects were fed marketed formula. Switching to a different marketed formula did not result in termination from the Term Formula phase.

		o de la composição de l	ocidel oy-Regimen P-value	0.87
			p.value	0.17
	hase	Study	p-value	0.00
	tudy formula p	Comparison	ייי אין רפי	0.967 0.998 0.998
Table 3	Weight Growth Rate During Study Formula Phase	Comparison	Control	Control vs DHA+ARA
			 	===
		Least Square S Mean	30.7	33.3 34.7
		c	09	3 %
		Regimen	Control	DIIA+ARA

Table 4

Weight at 40, 48, and 57 Weeks Post-Conceptual Age

eeks									
onceptual	_		. Least Square	Standard		Comparison	Study	Candar	· · · · · · · · · · · · · · · · · · ·
Age .	Regimen	c	Kean	Error	Comparison	p-value*	p-value	p.value	uender-by-Regimen p-value
07	Control	55	3075.3	67.9	Control vs DHA	0.388	0 50	3/ 0	;
	DIIA	24	3051.4	8.99	Control vs DHA+ARA	0.931	,	7.	1.00
	DIIA+ARA	29	3198.2	65.9	HM vs DHA	0.000			
	¥	8	3437.7	9.09	HM VS DHA+ARA	0.001			
	.=				IIM vs Control	0.000			
48	Control	53	4711.0	9.76	Control vs DHA	0.360	0.58	, ,	ć
	DIIA	51	4663.8	97.3	Control vs DHA+ARA	0.995)	2	0.29
	DIIA+ARA	. 57	5039.1	93.0	HH VS DIIA	0.000			
	¥	9	5181.5	85.9	HM VS DHA+ARA	0.114		٠	
	5				HH vs Control	0.000			
23	Control	14	6045.4	139.5	Control vs DHA	0.371	0.58	0 00	6.6
	DHA	65	5987.2	137.6	Control vs DNA+ARA	0.940	•	<u>;</u>	0.33
	DIIA+ARA	55	6312.9	127.9	HH VS DHA	0.005			
	Ξ	92	0.5079	126.7	IIM VS DIIA+ARA	0.278			
					IIN vs Control	0.014			

* One-sided test of the null hypothesis: Test Mean >= Control Mean

Table 5
Post-hoc Analysis of Weight

Time	Comparison	Two-sided p-value
Weight Gain During Study Formula Phase	C vs. DHA C vs. DHA+ARA DHA vs. DHA+ARA	0.067 0.004 0.30
Weight at 40 Weeks pca	C vs. DHA C vs. DHA+ARA DHA vs. DHA+ARA HM vs. DHA HM vs. DHA+ARA HM vs. C	0.78 0.14 0.074 <0.001 0.002 <0.001
Weight at 48 Weeks pca	C vs. DHA C vs. DHA+ARA DHA vs. DHA+ARA HM vs. DHA HM vs. DHA+ARA HM vs. C	0.72 0.011 0.004 <0.001 0.23 <0.001
Weight at 57 Weeks pca	C vs. DHA C vs. DHA+ARA DHA vs. DHA+ARA HM vs. DHA HM vs. DHA+ARA HM vs. C	0.74 0.12 0.057 0.010 0.56 0.028

Table 6

Length at 40, 48, and 57 Weeks Post-Conceptual Age

Gender-by-Regimen P-value 0.52 0.84 Gender P-value 0.05 Study p-value 0.03 0.00 0.00 Pairwise p-value 0.242 0.233 0.000 0.000 0.000 0.615 0.236 0.000 0.006 0.000 0.824 0.079 0.000 0.000 0.000 Control vs DHA
Control vs DHA+ARA
IH vs DHA
HH vs DIIA+ARA
Control vs HM
DHA vs DIIA+ARA Control vs DHA Control vs DHA+ARA Control vs DHA Control vs DHA+ARA HM vs DNA+ARA Control vs IIM DIIA vs DIIA+ARA HM VS DHA HM VS DHA+ARA Control VS HM DHA VS DHA+ARA Pairwise Comparison HM vs DHA Regimen p-value 0.000 0.000 0.000 Standard Least Square 48.4 47.8 49.0 50.6 54.7 54.6 55.5 57.4 60.7 60.5 61.3 62.4 52 54 58 83 83 53 52 57 81 22.22 Control DHA DHA+ARA HM Control DIIA DIIA+ARA IIM Control DHA DHA+ARA HM Regimen Weeks Post-Conceptual Age 70 48 27

Table 7

Head Circumference at 40, 48, and 57 Heeks Post-Conceptual Age

Gender-by-Regimen p-value	0.38	1.00	0.85
Gender p-value	0.00	0.00	0.00
Study p-value	0.91	0.81	79.0
Pairwise p-value	0.931 0.900 0.000 0.000 0.000 0.829		
Pairwise Comparison	Control vs DHA+ARA Econtrol vs DHA+ARA HH vs DHA HM vs DHA+ARA Control. vs HH DHA vs DHA+ARA		
Regimen p·value	000.0	0.983	0.689
Standard Error	2.00	0.2	0.2 0.2 0.2 0.2
Least Square Hean	35.4 35.4 35.5 36.5	39.1 39.0 39.0 39.0	41.9 41.6 41.7
c	53 58 85	52 51 56 81	53
	Control DHA DHA+ARA HM	Control DHA DHA+ARA HH	Control DHA DHA+ARA
Heeks Post∙Conceptual Age	. 07	87	25

Attorney Docket: 19400/09003 (MJ729)

Table 8 Visual Acuity at 48 and 57 Weeks Post-Conceptual Age

Study p-value	0.000	0.000
Pairwise p-value		0.697 0.071 0.042 0.000 0.113
Pairwise Comparison		Control vs DHA Control vs DHA+ARA HM vs DHA HM vs DHA+ARA Control vs HM
Regimen p-value	0.950	0.004
Standard Error (octaves)	. 0.10 0.10 0.09 0.09	0.08 0.08 0.07 0.07
Least Square Hean (log base2 cycles/deg)	0.76 0.85 0.78 0.81	1.79 1.75 1.61 1.94
Geometric mean (cycles/deg)	1.72 1.80 1.72 1.75	3.47
c	51 50 57 81	46 47 77 77
Regimen	Control DIIA DIIA+ARA IIH	Control DHA DHA+ARA HM
Veeks Post-Conceptual Age	8 7	57

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Pairwise p-value 0.196 0.010 0.176 Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA Pairwise Comparison Regimen p-value 0.762 0.559 0,165 0.884 0.441 0.243 0.679 0.830 0.034 Red Blood Cell Phosphatidylcholine Fatty Acids Hedian 0.036 0.030 0.031 0.599 0.686 0.656 0.021 0.016 0.018 36.594 35.578 35.987 0.845 0.976 0.931 11.468 11.201 11.174 17.308 16.935 16.988 18.952 19.603 18.824 Standard Error 0.019 0.013 0.009 0.036 0.031 0.031 0.009 0.005 0.006 0.540 0.462 0.445 0.049 0.243 0.238 0.192 0.298 0.391 0.271 0.525 0.505 0.466 Table 9 Arithmetic Hean 0.623 0.663 0.661 0.045 0.026 0.035 36.706 36.363 36.877 0.940 0.981 1.094 17.053 17.219 17.256 18.614 18.631 18.573 52 58 61 52 58 61 Regimen Control DHA DHA+ARA Control DHA DHA+ARA DHA DHA+ÀRA Control DHA DHA+ARA Control Control DHA DHA+ARA Fatty Acid 18:3n6 12:0 14:0 14:1 16:0 18:0 16:1 18:1 18:2 Study Form Initiation Study form Initiation

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		Pairwise p-value									
	•	Pairwise Comparison									
	Ø	Regimen p-value	0.647	0.234	0.723	0.290	0.673	0.507	0.819	0.155	0.911
	Fatty Acid	Median	0.224 0.236 0.188	0.246 0.246 0.216	0.262 0.281 0.269	0.000 0.017 0.008	0.632 0.640 0.614	2.096 2.296 2.135	8.124 7.876 8.207	0.105 0.130 0.139	0.298 0.302 0.329
Table 9	idylcholine	Standard Error	0.050 0.035 0.037	0.033	0.020 0.015 0.011	0.003	0.025 0.025 0.021	0.098 0.080 0.074	0.262 0.347 0.310	0.010 0.010 0.010	0.057 0.015 0.015
Tab	Red Blood Cell Phosphatidylcholine Fatty Acids	Arithmetic Mean	0.399 0.337 0.310	0.315 0.257 0.233	0.287 0.287 0.268	0.017 0.025 0.017	0.632 0.628 0.602	2.144 2.208 2.218	7.657 8.164 8.090	0.106 0.127 0.126	0.351 0.322 0.321
	B100d (c	52 58 61								
	Red	Regimen	Control DHA DHA+ARA								
		fatty Acid	20:0	18:3 <i>n</i> 3	20:1	18:4	20:2n6	20:3n6	20:4n6	22:1	20:5n3
J- 1	-	Time	Study Form Initiation								

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				Tab	Table 9				
		Red	81 ood	Red Blood Cell Phosphatidylcholine Fatty Acids	idylchol ine	fatty Acid	Ś		
T i me	Fatty Acid	Regimen	c	Arithmetic Mean	Standard Error	Median	Regimen p-value	Pairwise Comparison	Pairwise p-value
Study form Initiation	32:4n6	Control DHA DHA+ARA	52 58 61	0.578 0.493 0.443	0.144 0.030 0.021	0.423 0.481 0.425	0.331		
Study Form Initiation.	24:1	Control DHA DHA+ARA	52 58 61	0.208 0.115 0.180	0.054 0.019 0.056	0.075 0.084 0.096	0.665		
Study form Initiation	22:5n6	Control DHA DHA+ARA	52 58 61	0.266 0.259 0.265	0.020 0.017 0.018	0.232 0.239 0.256	0.923		
Study form Initiation	22:4n3	Control DHA DHA+ARA	52 58 61	0.000	0.000 0.001 0.001	0.000	0.199		
Study Form Initiation	. 22:5n3	Control DHA DHA+ARA	52 58 61	0.213 0.215 0.203	0.019 0.013 0.010	0.203 0.195 0.193	0.885		
Study Form Initiation	22:6n3	Control DHA DHA+ARA	52 58 61	0.984 1.075 1.006	0.051 0.053 0.050	1.000 1.034 0.970	0.858		

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		Pairwise p-value					0.118 0.003 0.152			0.600	
		Pairwise Comparison					Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA			Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA	
		Regimen p-value	0.843	0.834	0.155	0.767	0.013	0.886	0.686	0.001	0.527
	fatty Acids	Hedian	0.035 0.031 0.032	0.806	0.033 0.015 0.018	34.798 34.841 33.890	0.526 0.475 0.472	14.197 13.867 14.108	14.291 13.998 14.218	21.506 22.517 20.662	0.074 0.076 0.066
Table 9	idylcholine	Standard Error	0.026 0.042 0.012	0.039 0.035 0.036	0.008 0.009 0.007	0.512 0.595 0.584	0.026 0.042 0.029	0.261 0.237 0.253	0.277 0.272 0.380	0.340 0.457 0.337	0.006
Tab	Red Blood Cell Phosphatidylcholine Fatty Acids	Arithmetic Mean	0.100	0.808 0.781 0.755	0.047 0.036 0.036	35.837 35.560 35.069	0.566 0.594 0.526	13.972 14.065 14.341	14.456 14.116 14.344	21.673 22.045 19.899	0.080 0.088 0.087
	Blood C	· c	55 59 59	53 56 59	53 56 59	55 53	53 59 59	53 56 59	53 59 59	53 26 26	55 55
	Red	Regimen	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DIIA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA
		Fatty Acid	12:0	14:0	14:1	16:0	16:1	18:0	18:1	18:2	18:3n6
	13 to - **	1 ime	Study form Termination	Study Form Termination	Study Form Termination	Study Form Termination	Study Form Termination	Study form Termination	Study Form Termination	Study Form Termination	Study Form Termination

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Table 9

Red Blood Cell Phosphatidylcholine Fatty Acids

Pairwise P-value		0.503 0.068 0.011	:				0.097 0.000 0.000		0.004 0.108 0.000
Pairwise Comparison		Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA					Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA		Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA
Regimen p-value	0.424	0.031	0.149	0.672	0.051	0.208	0.000	0.946	0.000
Hedian	0.392 0.281 0.251	0.283	0.302 0.283 0.283	0.015 0.018 0.008	0.910 0.873 0.821	2.091 2.043 1.904	6.029 5.892 8.891	0.125 0.114 0.104	0.189 0.233 0.169
Standard Error	0.050 0.053 0.049	0.020 0.030 0.009	0.014 0.013 0.013	0.004 0.003 0.002	0.026 0.023 0.022	0.073 0.070 0.064	0.240 0.220 0.255	0.010 0.009 0.011	0.022 0.012 0.014
Arithmetic Mean	0.504 0.472 0.430	0.321 0.335 0.273	0.318 0.300 0.307	0.022 0.022 0.014	0.893 0.880 0.824	2.032 2.017 1.908	6.046 5:774 8.465	0.117 0.110 0.115	0.214 0.246 0.186
. =	55.55	59 82	28 82	53 56 59	25 53	25 22	23 26 26	25 55 59	288
Regimen	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DHA÷ARA	Control DHA DHA+ARA	Control DHA DNA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA
fatty Acid	20:0	18:3n3	20:1	18:4	20:2n6	20;3n6	20:4n6	22:1	20:5n3
T i me	Study form Termination	Study form Termination:	Study Form Termination	Study Form Termination	Study Form Termination				

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Pairwise p-value

Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA Pairwise Comparison Regimen p-value 0.000 900.0 0.359 0.221 0.093 0.303 Red Blood Cell Phosphatidylcholine Fatty Acids 0.812 1.352 1.259 0.289 0.260 0.255 Median 0.062 0.086 0.089 0.163 0.133 0.165 0.000 0.390 0.426 0.487 Standard Error 0.019 0.026 0.013 0.072 0.063 0.049 0.039 0.036 0.040 0.013 0.011 0.009 0.001 0.001 0.002 0.048 0.061 0.027 Table 9 Arithmetic Hean 0.895 1.380 1.244 0.306 0.293 0.265 0.001 0.001 0.003 0.484 0.489 0.496 0.127 0.143 0.177 0.181 0.145 0.172 288 Control DHA DHA+ARA DHA DHA+ARA Control DHA DHA+ARA DHA DHA+ARA Control DHA DHA+ARA DHA DHA+ARA Control Control Regimen Control 22:6n3 22:5n3 22:5n6 22:4n3 22:4n6 fatty Acid 24:1 Study Form Termination lime

0.005 0.895 0.006 0.000 0.000 0.141

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		Pairwise p-value				0.527 0.593 0.000 0.000 0.000	0.524 0.467 0.000 0.000 0.000 0.183
		Pairwise Comparison				Control vs DHA Control vs DHA+ARA HM vs DHA HM vs DHA+ARA Control vs HM DHA vs DHA+ARA	Control vs DHA Control vs DHA+ARA IIH vs DHA HH vs DHAARA Control vs HH
	Acids	Regimen p-value	0.729	0.943	0.448	0.000	0.000
	line fatty	Median	0.026 0.016 0.021 0.020	0.331 0.324 0.328 0.335	0.013 0.011 0.015 0.020	34.319 34.473 34.165 32.228	0.338 0.352 0.368 0.473
Table 9	phatidylcho	Standard Error	0.005 0.006 0.004 0.016	0.039 0.032 0.024 0.026	0.006 0.007 0.006 0.003	0.577 0.689 0.506 0.506	0.043 0.023 0.024 0.020
	Red Blood Cell Phosphatidylcholine Fatty Acids	Arithmetic Meàn	0.032 0.028 0.026 0.059	0.402 0.353 0.353 0.381	50.0 9.0.0 9.0.0 9.0.0	34.627 35.272 34.802 33.037	0.435 0.380 0.395 0.507
	Red BI	c	37 38 38 56	32 38 38 56	32 38 38 56	37 38 38 56	37 38 38 56
		Regimen	Control DHA DHA+ARA HH	Control DHA DHA+ARA HH	Control DHA DHA+ARA HM	Control DHA DHA+ARA HM	Control DHA DHA+ARA HM
		Fatty Acid	12:0	14:0	14:1 	9:0	16:1
	·	Time	48 Weeks PCA	48 Weeks PCA	48 Veeks PCA	48 Weeks PCA	48 Weeks PCA

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		Pairwise p·value	0.760 0.889 0.000 0.000 0.000		0.840 0.527 0.000 0.000 0.000	0.950 0.774 0.004 0.001 0.003	
		Pairwise Comparison	Control vs DHA Control vs DHA+ARA HH vs DHA HH vs DHA+ARA Control vs HM DHA vs DHA+ARA		Control vs DHA Control vs DHA+ARA HM vs DHA HM vs DHA+ARA Control vs HM DHA vs DHA+ARA	Control vs DHA Control vs DHA+ARA HH vs DHA HH vs DHA+ARA Control vs HH DHA vs DHA+ARA	
	Acids	Regimen p-value	0.000	0.256	0.000	0.005	0.785
	line Fatty	Median	12.759 12.786 12.793 14.729	18.636 18.492 18.227 18.727	23.552 23.717 23.839 18.482	0.061 0.067 0.062 0.039	0.197 0.206 0.172 0.215
Table 9	phatidylcho	Standard Error	0.313 0.249 0.235 0.287	0.453 0.289 0.305	0.518 0.516 0.422 0.344	0.008 0.005 0.006 0.004	0.075 0.061 0.061 0.044
	Red Blood Cell Phosphatidylcholine Fatty Acids	Arithmetic Hean	13.016 12.944 12.804 14.583	17.894 17.766 17.850 18.662	23.469 23.538 23.738 18.650	0.071 0.069 0.069 0.042	0.348 0.339 0.304 0.409
	Red B	c	37 38 38 56	37 32 38 56	33 38 38 38	28 33 3	32 33 38 38 39
		Regimen	Control DHA DHA+ARA HH	Control DHA DHA+ARA HM	Control DHA DHA+ARA . HH	Control DHA DHA+ARA HH	Control DHA DHA+ARA HH
. =		Fatty	18:0	18:1	18:2	18:3n6	50:0
		lime	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA

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		Pairwise p-value	0.812 0.918 0.001 0.002 0.001	0.579 0.588 0.001 0.001 0.000 0.974	0.822 0.161 0.039 0.001 0.054		0.610 0.735 0.000 0.000 0.000
	-	Pairwise Comparison	Control vs DHA+ARA Control vs DHA+ARA HH vs DHA HH vs DHA+ARA Control vs HH DHA vs DHA+ARA	Control vs DHA+ARA Control vs DHA+ARA HH vs DHA HH vs DHA+ARA Control vs HH DHA vs DHA+ARA	Control vs DHA+ARA Control vs DHA+ARA HM vs DHA IIM vs DHA+ARA Control vs HH DHA vs DHA+ARA		Control vs DHA Control vs DHA+ARA HH vs DHA HH vs DHA+ARA Control vs HH DHA vs DHA+ARA
;	Acids	Regimen p-value	0.001	0.000	0.010	0.629	0:000
:	line Fatty	Median	0.182 0.182 0.190 0.120	0.420 0.435 0.375 0.309	0.000 0.000 0.000 0.015	0.537 0.543 0.550 0.531	1.741 1.684 1.717 2.166
Table 9	phatidylcho	Standard Error	0.019 0.015 0.010 0.022	0.019 0.025 0.016 0.014	0.005 0.004 0.005 0.004	0.023 0.032 0.053 0.014	0.086 0.073 0.090 0.086
; ;	Red Blood Cell Phosphatidylcholine Fatty Acids	Arithmetic Mean	0.222 0.211 0.203 0.182	0.416 0.406 0.382 0.311	0.018 0.016 0.007 0.024	0.543 0.557 0.636 0.560	1.709 1.702 1.844 2.265
-	Red B	د `	37 32 38 56	37 32 38 56	28 32 32	32 33 35 35 35 35 35 35 35 35 35 35 35 35	32 38 35 56 56 56 56 56 56 56 56 56 56 56 56 56
		Regimen	Control DHA DHA+ARA HM	Control DHA DHA+ARA HH	Control DHA DHA+ARA HM	Control DHA DHA+ARA HM	Control DHA DHA+ARA HM
		Fatty Acid	18:303	20:1	18:4	20:2n6	20:3n6
		T ime	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA

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Pairwise p-value 0.633 0.086 0.000 0.000 0.000 0.508 0.805 0.000 0.000 0.000 0.337 0.247 0.000 0.000 0.000 Control vs DHA
Control vs DHA+ARA
HH vs DHA
HN vs DHA+ARA
Control vs HH
DHA vs DHA+ARA Control vs DHA Control vs DHA+ARA Control vs DHA Control vs DHA+ARA HM vs DHA HM vs DHA+ARA Control vs HM DHA vs DHA+ARA HM VS DHA HM VS DHA+ARA Control VS HM DHA VS DHA+ARA Comparison Pairwise Regimen p-value 0.000 0.000 0.000 0.244 0.664 Red Blood Cell Phosphatidylcholine Fatty Acids Hedian 4.736 4.499 4.746 7.666 0.131 0.118 0.105 0.104 0.077 0.083 0.078 0.123 0.373 0.417 0.384 0.377 0.112 0.116 0.108 0.079 Standard Error Table 9 0.070 0.062 0.055 0.020 0.036 0.014 0.024 0.030 0.015 0.006 0.009 0.009 0.059 0.029 0.054 0.022 0.255 0.196 0.185 0.250 Arithmetic Mean 0.102 0.084 0.099 0.138 0.426 0.382 0.440 0.406 0.247 0.210 0.179 0.115 0.166 0.116 0.131 0.160 4.738 4.475 4.550 7.408 37 38 38 56 32 38 38 56 32 33 36 36 36 28 32 32 32 38 38 56 DHA DHA+ARA HM Control DHA DHA+ARA HH Control DHA DHA+ARA HM Control DHA DHA+ARA HM DHA+ARA HM Control Regimen Control 20:503 22:4n6 20:4n6 Fatty Acid 24:1 22:1 48 Weeks PCA Time

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Pairwise p-value 0.598 0.759 0.000 0.000 0.000 0.505 0.647 0.000 0.001 0.000 0.111 0.052 0.000 0.000 0.000 Control vs DHA Control vs DHA+ARA HH vs DHA Control vs DHA Control vs DHA+ARA Control vs DHA Control vs DHA+ARA HH vs DHA IIN vs DHA+ARA Control vs HM DHA vs DHA+ARA HM vs DHA+ARA Control vs HM DHA vs DHA+ARA HM vs DHA HM vs DHA+ARA Control vs HM DHA vs DHA+ARA Pairwise Comparison Regimen p·value 0.000 0.000 0.000 1.000 Red Blood Cell Phosphatidylcholine Fatty Acids 0.212 0.186 0.198 0.265 Standard Error 0.029 0.017 0.026 0.018 0.047 0.048 0.043 0.043 Table 9 0.016 0.012 0.022 0.016 Arithmetic Mean 0.595 0.685 0.662 1.475 0.000 0.210 0.189 0.231 0.264 2883 38 32 32 32 32 32 32 37 38 38 56 38 38 Control DHA DHA+ARA HM Control DHA DHA+ARA HM Control DHA DHA+ARA HM Regimen Control DHA DHA+ARA HM 22:6n3 22:5n3 22:4n3 22:5n6 Fatty 48 Weeks PCA 48 Weeks PCA 48 Weeks PCA 48 Weeks PCA

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Pairwise p-value 0.373 Control vs DHA+ARA Control vs DHA+ARA DHA vs DHA+ARA Pairwise Comparison Regimen p-value 0.546 0.792 0.181 . 296.0 0.337 0.142 0.412 0.773 0.000 Red Blood Cell Phosphatidylethanolamine Fatty Acids 16.698 16.308 16.001 0.145 0.152 0.169 Median 0.698 0.746 0.837 8.469 8.308 7.904 6.682 6.346 5.682 0.022 0.033 0.039 0.220 0.206 0.246 0.032 0.028 0.050 17.945 19.295 19.035 Standard Error 0.018 0.019 0.016 0.038 0.025 0.021 0.035 0.034 0.035 0.329 0.227 0.215 0.301 0.326 0.375 0.015 0.013 0.010 0.015 0.012 0.009 0.736 0.622 0.451 Table 10 Arithmetic Hean 20.021 19.847 19.796 0.731 0.769 0.836 16.450 16.208 16.415 6.615 6.336 6.175 8.857 8.434 8.201 0.307 0.278 0.277 0.080 0.061 0.062 0.069 0.075 0.063 52 54 2 Control DHA DHA+ARA Control DHA DHA+ARA Control **DHA+ARA** Control DHA DHA+ARA Control DHA DHA+ARA Control DHA DHA+ARA Control DHA DHA+ARA Regimen Control Control DHA+ARA 18:3n6 fatty Acid 18:0 18:1 18:2 12:0 14:0 16:0 16:1 14:1 Study form Initiation Jime

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		Pairwise D-value									
		Pairwise Comparison					·				
	cids	Regimen p•value	0.151	0.641	0.395	0.371	0.706	660.0	0.353	0.572	0.997
	ine fatty A	Median	0.291 0.244 0.186	0.261 0.249 0.225	0.517 0.555 0.544	0.000 0.025 0.021	0.480 0.437 0.427	1.829 1.820 1.911	26.820 27.376 27.708	0.138 0.151 0.141	0.357 0.370 0.335
10	/lethanolami	Standard Error	0.043 0.030 0.024	0.023 0.018 0.016	0.036 0.034 0.027	0.005 0.004 0.007	0.023 0.024 0.028	0.072 0.077 0.064	0.618 0.611 0.645	0.017 0.015 0.017	0.024 0.024 0.022
Table 10	Red Blood Cell Phosphatidylethanolamine Fatty Acids	Arithmetic Mean	0.372 0.314 0.259	0.305 0.269 0.257	0.573 0.615 0.571	0.025 0.031 0.030	0.479	1.843 1.965 1.973	25.817 26.475 26.747	0.150 0.167 0.168	0.378 0.384 0.366
	lood Ce	<u>د</u>	52 57 61	52 57 61	52 57 61						
	Red B	Regimen	Control DHA DHA+ARA	Control DHA DHA+ARA	Control DHA DHA+ARA						
		Fatty Acid	20:0	18:3n3	20:1	18:4	20:2n6	20:3n6	20:4n6	22:1	20:5n3
		. Time	Study Form Initiation	Study Form Initiation	Study Form Initiation						

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Pairwise p-value Pairwise Comparison 0.375 0.875 0.068 0.555 0.195 0.257 Red Blood Cell Phosphatidylethanolamine fatty Acids 0.000 Hedian 1.782 1.857 1.775 0.041 0.031 0.047 0.083 0.070 0.075 0.001 0.001 0.002 0.028 0.009 0.010 Table 10 Arithmetic Kean 0.100 0.059 0.072 1.757 1.809 1.851 0.001 0.001 0.005 52 57 61 Control DHA DHA+ARA 22:513 22:6n3 22:5n6 22:4n3 22:4n6 fatty Acid 24:1 Study Form Initiation Study form Initiation

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Pairwise p-value 0.130 0.006 0.219 0.908 0.000 0.000 Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA Pairwise Comparison Regimen p-value 0.630 0.000 0.782 0.592 0.560 0.604 0.024 0.333 0.160 Red Blood Cell Phosphatidylethanolamine Fatty Acids 9.406 8.818 8.697 0.033 0.036 0.035 0.279 0.265 0.256 0.041 0.000 0.043 17.617 17.556 17.568 0.476 0.509 0.555 14.695 14.927 14.499 0.163 0.157 0.161 Standard Error 0.034 0.045 0.049 0.192 0.207 0.141 0.018 0.019 0.012 0.031 0.039 0.030 0.020 0.013 0.011 0.673 0.614 0.467 0.266 0.208 0.242 0.437 0.299 0.330 0.012 0.017 0.018 Table 10 Arithmetic 0.511 0.579 0.618 0.360 0.380 0.348 0.086 0.066 0.066 19.326 19.062 18.357 9.614 9.173 8.961 14.763 15.177 14.814 0.093 0.093 0.067 58.53 28 23 23 Control DHA DHA+ARA DHA+ARA Regimen DHA DHA+ARA DHA DHA+ARA DHA DHA+ARA Control Control DHA Control DHA DHA+ARA Control Control Control DHA+ARA DIIA+ARA Control Control 18:3n6 Fatty Acid 18:2 18:0 18:1 12:0 14:0 14:1 16:0 16:1 Study Form Termination I ime

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				Table 10	10				
		. Red B	ood Ce	Red Blood Cell Phosphatidylethanolamine Fatty Acids	ylethanolami	ne Fatty A	cids		
Time	Fatty Acid	Regimen	<u>.</u>	Arithmetic Mean	Standard Error	Median	Regimen p-value	Pairwise Comparison	Pairwise p-value
tudy form lermination	20:0	Control DHA DHA+ARA	53 58	0.404 0.336 0.288	0.044	0.278 0.208 0.208	0.146		
itudy Form Termination	18:3n3	Control DHA DHA+ARA	55 58 58	0.382 0.368 0.329	0.017 0.016 0.015	0.364 0.354 0.305	0.134		
study Form Termination	20:1	Control DIIA DIIA+ARA	22 23	0.553 0.579 0.507	0.029 0.028 0.025	0.526 0.537 0.483	0.164		
Study Form Termination	18:4	Control DHA DHA+ARA	888	0.042 0.026 0.022	0.010 0.005 0.004	0.018 0.019 0.000	0.108		
Study Form Termination	20:2n6	Control DHA DHA+ARA	ននន	0.754 0.774 0.654	0.029 0.030 0.026	0.765 0.750 0.663	0.068		
study Form Termination	20:3n6	Control DHA DHA+ARA	8 23	2.253 2.295 2.066	0.111 0.094 0.073	2.073 2.206 1.992	0.203		
Study Form Termination	20:4116	Control DHA DHA+ARA	53 58	24.279 23.464 26.760	0.527 0.520 0.437	25.132 24.038 27.372	0.000	Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA	0.119 0.000 0.000
Study Form Termination	22:1	Control DHA DHA+ARA	53 58	0.149 0.176 0.146	0.019 0.016 0.012	0.122 0.169 0.130	0.229		
Study form Termination	20:5n3	Control DHA DHA+ARA	53 58	0.519	0.020 0.025 0.015	0.493 0.575 0.415	00000	Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA	0.286

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-				Tabi	Table 10				
		Red B	l ood C	Red Blood Cell Phosphatidylethanolamine Fatty Acids	ylethanolami	ine Fatty A	cids		
Time	fatty Acid	Regimen	ċ	Arithmetic Mean	Standard Error	Median	Regimen p-value	Pairwise Comparison	Pairwise D-value
Study form Termination	22:4n6	Control DHA DHA+ARA	53 58.	7.309 7.135 7.592	0.208 0.154 0.155	7.656 6.885 7.635	0.007	Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA	0.025
Study form Termination	24:1	Control DHA DHA+ARA	\$ 23	0.092 0.056 0.062	0.023	0.038	0.294		
Study Form Termination	22:5n6	Control DHA DHA+ARA	28 23	1.444 1.231 1.347	0.064 0.034 0.040	1.423	0.010	Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA	0.003
Study form Termination	22:4n3	Control DHA DHA+ARA	55.53	0.000	0.000	0.000	0.137	·	
Study form Termination	22:5n3	Control DHA DHA+ARA	53 58	2.694 2.334 2.237	0.110 0.091 0.069	2.839 2.400 2.269	0.003	Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA	0.00¢ 0.002 0.943
Study Form Termination	22:6n3	Control DHA DHA+ARA	55 53	4.798 6.762 6.389	0.151 0.183 0.150	4.815 7.043 6.498	0.000	Control vs DHA Control vs DHA+ARA DHA vs DHA+ARA	0.000

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	Pairwise p-value					0.601 0.524 0.000 0.000 0.001
	Pairwise Comparison					Control vs DIIA Control vs DHA+ARA HM vs DHA HH vs DHA+ARA Control vs HH DHA vs DHA+ARA
ty Acids	Regimen p·value	0.587	0.598	0.092	0.177	0.000
Jamine Fati	Median	0.024 0.019 0.018 0.023	0.169 0.162 0.188 0.210	0.037 0.000 0.044 0.021	16.314 15.692 16.997 17.607	0.349 0.336 0.376 0.562
Table 10 atidylethanc	Standard Error	0.019 0.016 0.014 0.011	0.030 0.041 0.025 0.016	0.017 0.017 0.019 0.011	0.595 0.729 0.538 0.395	0.050 0.035 0.022 0.027
Table 10 Red Blood Cell Phosphatidylethanolamine Fatty Acids	Arithmetic Mean	0.053 0.054 0.047 0.045	0.243 0.251 0.235 0.236	0.080 0.055 0.078 0.053	17.319 17.101 17.225 18.138	0.440 0.390 0.390 0.390
ed Bloc	c	37 32 38 56	37 32 38 56	37 38 38 56	37 38 38 56	37 32 38 56
<u>~</u>	Regimen	Control DHA DHA+ARA HM	Control DHA DHA+ARA HM	Control DHA DHA+ARA HM	Control OHA DHA+ARA HM	Control DHA DHA+ARA HH
	fatty Acid	:= 5 <u>7</u> ::	14:0	14:1	16:0	19:1
	T i me	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA

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Pairwise p-value 0.347 0.483 0.020 0.000 0.001 0.108 0.401 0.234 0.067 0.118 0.005 0.758 Control vs DHA Control vs DHA+ARA IM vs DHA+ARA Control vs HM DHA vs DHA+ARA HN vs DHA+ARA Control vs HM DHA vs DHA+ARA Control vs HM DHA vs DHA+ARA HN VS DHA HH VS DHA+ARA Pairwise Comparison HM vs DHA HN VS DHA HH VS DHA Regimen p-value Red Blood Cell Phosphatidylethanolamine Fatty Acids 0.000 0.038 0.00 0.050 0.728 Median 7.174 7.552 7.173 8.409 19.410 19.534 19.433 18.141 9.267 8.696 8.840 6.027 0.182 0.171 0.158 0.158 0.146 0.145 0.125 0.240 Standard Error Table 10 0.327 0.293 0.270 0.230 0.368 0.421 0.332 0.278 0.261 0.210 0.216 0.193 0.020 0.031 0.021 0.012 0.058 0.042 0.037 0.031 Arithmetic Hean 7.935 7.962 7.443 8.754 19.438 19.066 19.302 18.469 9.328 8.867 9.257 6.291 0.263 0.262 0.212 0.295 32 33 38 38 38 38 28 33 32 38 33 26 38 26 38 38 32 38 26 28 29 28833 Control DHA DHA+ARA HN Control DHA DHA+ARA HM DHA DHA+ARA HM Control DHA DHA+ARA HM Regimen Control DHA DHA+ARA HN Control 18:3n6 fatty Acid 18:0 18:1 18:2 20:02 48 Weeks PCA Time

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Pairwise p-value 0.559 0.848 0.008 0.002 0.001 0.689 0.339 0.512 0.000 0.000 0.000 0.543 0.532 0.000 0.000 0.000 0.896 0.935 0.015 0.006 0.007 0.835 Control vs DHA+ARA HH vs DHA HH vs DHA+ARA Control vs HH DHA vs DHA+ARA Control vs DHA Control vs DHA+ARA Control vs DHA Control vs DHA+ARA Control vs DHA Control vs DHA+ARA HM vs DHA HM vs DHA+ARA Control vs HM DHA vs DHA+ARA HM VS DHA HM VS DHA+ARA Control VS HM DHA VS DHA+ARA Control vs DHA HN VS DHA HN VS DHA+ARA Control VS HM DHA VS DHA+ARA Pairwise Comparison Regimen p-value Red Blood Cell Phosphatidylethanolamine fatty Acids 0.000 0.000 0.001 0.057 0.012 Median 0.225 0.262 0.245 0.169 0.648 0.782 0.738 0.492 0.003 0.000 0.000 0.019 0.698 0.684 0.689 0.412 1.999 2.045 2.132 1.637 Standard Error Table 10 0.025 0.017 0.015 0.020 0.031 0.032 0.188 0.024 0.005 0.005 0.006 0.006 0.035 0.026 0.032 0.016 0.099 0.100 0.114 0.053 Arithmetic Kean 0.291 0.270 0.265 0.226 0.672 0.668 0.715 0.444 0.715 0.772 0.936 0.533 0.017 0.017 0.023 0.027 2.138 2.165 2.172 1.715 28 32 32 32 32 37 38 38 56 28 32 22 2883 28 32 32 Control DHA DHA+ARA HM Control DHA DHA+ARA HM Control DHA DHA+ARA HM DHA DHA+ARA HM Regimen Control DHA DHA+ARA HH Control 18:3n3 20:2n6 20:3n6 fatty Acid 20:1 48 Weeks PCA T i me

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		Pairwise p-value				0.612 0.416 0.000 0.013 0.001	
		Pairwise Comparison			·	Control vs DHA Control vs DHA+ARA IIH vs DHA HH vs DHA+RRA Control vs IIH DHA vs DIA+ARA	
	tty Acids	Regimen p-value	0.950	0.121	265.0	0.001	0.943
	olamine Fa	Median	24.774 25.206 25.122 25.122	0.172 0.188 0.133 0.134	0.368 0.377 0.347 0.360	8.761 9.132 8.472 7.618	0.035 0.034 0.036 0.027
Table 10	hatidylethar	Standard Error	0.536 0.491 0.429 0.384	0.016 0.022 0.022 0.013	0.026 0.015 0.011 0.016	0.267 0.250 0.188 0.203	0.016 0.009 0.008 0.016
	Red Blood Cell Phosphatidylethanolamine Fatty Acids	Arithmetic Mean	24.508 24.428 24.788 24.625	0.168 0.189 0.154 0.148	0.382 0.369 0.347 0.384	8.580 8.791 8.576 7.727	0.067 0.049 0.046 0.062
	Red Blo	c	37 38 38 56	37 38 38 56	37 38 38 56	33 38 38 56	37 38 38 56
		Regimen	Control DHA DHA+ARA HM	Control DHA DHA+ARA HM	Control DHA DHA+ARA HH	Control OHA DHA+ARA HM	Control DHA DHA+ARA HM
		Fatty	20:4n6	22:1	20:5n3	22:406	24:1.
		ĭime	48 Weeks PCA	48 Weeks PCA	48 Weeks PCA	48 Neeks PCA	48 Weeks PCA

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Table 10

Pairwise P-value 0.977 0.997 0.000 0.000 0.000 0.884 0.148 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 Control vs DHA
Control vs DHA+ARA
HM vs DHA
HH vs DHA+ARA
Control vs HM Control vs DHA+ARA Control vs DHA Control vs DHA+ARA HH VS DHA+ARA Control vs HM DHA VS DHA+ARA Control vs DHA IIH VS DIIA HM VS DIIA+ARA Control VS HM DHA VS DHA+ARA Comparison HM vs DHA Pairwise Regimen p-value Red Blood Cell Phosphatidylethanolamine Fatty Acids 0.000 1.000 0.000 0.000 Median 1.414 1.359 1.889 0.000 0.000 0.000 0.000 2.681 2.630 2.443 1.978 3.013 4.079 3.721 7.341 Standard Error 0.066 0.057 0.054 0.056 0.000 0.000 0.000 0.001 0.092 0.086 0.066 0.065 Arithmetic Mean 0.000 0.000 0.000 0.001 2.567 2.561 2.436 1.942 1.401 1.353 1.364 1.883 3.196 4.143 3.801 7.283 28 32 32 28832 28 22 22 DHA DHA+ARA HM Control DHA DHA+ARA HM Control DHA DHA+ARA HM Control DHA DHA+ARA HH Regimen Control 22:5n6 22:4n3 22:5n3 22:613 fatty Acid 48 Weeks PCA 48 Weeks PCA 48 Weeks PCA 48 Weeks PCA Time

Table 11
Preterm Infant Complications

		Regimen		p-value*
	Control	DHA	DHA+ARA	1
Retinopathy of Prematurity Test Results Absent I II III Present, but not graded	34 (76%) 8 (18%) 2 (4%) 1 (2%)	44 (76%) 11 (19%) 2 (3%) 1 (2%)	41 (79%) 6 (12%) 4 (8%) 1 (2%)	0.91
Ultrasound Examination for Intraventricular Hemorrhage None Stage 1 Stage 2 Stage 3 Stage 4 Questionable	47 (81%) 6 (10%) 3 (5%) 1 (2%) 1 (2%)	52 (84%) 9 (15%) 1 (2%)	49 (80%) 7 (11%) 2 (3%) 1 (2%) 2 (3%)	0.78
Posthemorrhagic Hydrocephalus developed? No Yes	61 (98%) 1 (2%)	65 (98%) 1 (2%)	64 (97%) 2 (3%)	1.00

^{*}The statistical test was based on a dichotomous response: present or absent.

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Table 12
Serious Adverse Events Reported During Study Formula Phase

		Regimen		
Event	Control	DHA	DHA+ARA	p-value
Any Event	4 (6%)	3 (5%)	4 (6%)	0.93
Other Respiratory Conditions of Fetus and Newborn	2 (3%)	0	0 .	0.10
Other Infection Specific to the Perinatal Period	1 (2%)	0	0	0.32
Intraventricular Hemorrhage	0	0	1 (2%)	1.00
Other Specified Perinatal Disorders of Digestive System	0	1 (2%)	0	1.00
Convulsions in Newborn	1 (2%)	0	0	0.32
Feeding Problems in Newborn	0	1 (2%)	1 (2%)	1.00
Hernia	0	0	1 (2%)	1.00
Other	0	1 (2%)	1 (2%)	1.00

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Table 13
Serious Adverse Events Reported During the Term Formula Phase

		De	gimen	·	
Event		T			
	Control	+	DHA + ARA	HIM	p-value
Any Event	7 (13%)	9 (15%)	9 (15%)	1 (1%)	0.002 C VS D 0.79 C VS D+A 0.79 D VS D+A 1.00 C VS HM 0.006 D VS HM 0.001 D+A VS HM 0.001
Infectious Colitis, Enteritis, and Gastroenteritis	0	o	1 (2%)	0	0.67
Croup	0	o	1 (2%)	0	0.67
Bronchopneumonia, Organism Unspecified	2 (4%)	3 (5 1)	6 (10%)	o	0.013 C vs D 1.00 C vs D+A 0.27 D vs D+A 0.49 C vs HM 0.15 D vs HM 0.064 D+A vs HM 0.004
Asthma, Unspecified	1 (2%)	0	0	0	0.21
Esophageal Reflux	0	1 (2%)	.2 (3%)	0	0.23
Dyspepsia and Other Stomach Function Disorder	0	0	0	1 (1%)	1.0
Other Respiratory Conditions of Fetus and Newborn	1 (2%)	1 (2%)	3 (5%)	0	0.11
Convulsions	1 (2%)	0	0	0 .	0.21
udden Infant Death yndrome	1 (2%)	1. (2%)	0	0	0.34
lernia	2 (4%)	2 (3%)	0	0	0.11
ther	0	3 (5%)	2 (3%)	0	0.063

Appendix 1

Listing of Weights Included in the Statistical Analyses

49t_57 6816 56.6 6610 7470 57.3 49c 48 3731 3064 3575 3688 3745 3070 3070 3590 3620 40.1 2520 Growth Rate g/day 23.9 56.9 31.5 34.1 36.1 36.2 33.8 27.7 43.3 34.2 28.9 41.7 54.4 Wg t'9 Wgt8 Hgt7 2045 Ng t 6 1760 37.3 2340 2012 2425 1665 36.3 1870 1450 35.4 2045 33.0 1494 34.4 1851 1840 35.4 1566 2040 1230 32.6 1261 32.0 1855 32.6 1298 1775 32.1 1205 1630 33.4 975.0 32.3 1600 34.4 1810 1785 33.3 30.7 Weight (g) Age (weeks pca) Veight (g) Age (weeks pca) Veight (g) Age (weeks pca) Veight (g) Age (weeks pca) pca) Veight (g) Age (weeks pca) Weight (g) Age (weeks pca) Veight (g) Age (weeks pca) Weight (g) Age (weeks pca) pca) Weight (g) Age (weeks pca) Veight (g) Age (weeks pca) Age (weeks pca) Weight (g) Age (weeks p Weight (g) Age (weeks | Weight (g) Variable 9703-0304 9704-0303 9050-6696 9010-6696 9701-0303 9701-0304 9702-0302 9703-0302 9703-0308 9699-0302 9700-0301 9698-0304 9698-0301 Subject Control. Control Control Control Control Control Control Control Control Gender Hale Male Hale Hale Hale Male Hale Male Hale Male Male Hale Hale

subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

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Analyses
Statistical
in the
<u>≃</u> .
Included
Weights
ō
Listing
<i>:</i> .

Nat 1 Mat 2
1640 33.0
1588 35.0
1570
2130
1984 2135 34.7 35.6
1734 2005 33.1 34.0
1820 2215 32.9 34.4
1600 1850 34.1 35.1
1442 1644 32.7 33.7
1960 2205 33.7 34.7
1440 1660 31.7 32.7
1221 1245 31.7 31.9
1345 1456 34.1 35.1

ppendix 1

sting of Weights Included in the Statistical Analyses

Grouth Rate 10.0 2260 4535 10.0 2260 4535 10.0 2260 4776 6695 48.9 3085 4795 6695 48.9 3085 4795 6695 48.9 3085 4795 6695 48.9 3085 4795 6695 48.9 3085 4796 67036 48.0 37.0 48.3 2775 434 6022 48.3 2779 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.3 2774 48.1 57.0 48.0 57.0 49.0 2104 2276 2288 20.4 2805 3405 46.0 52.0 39.0 48.0 5280 1.2 2019 2104 2276 2288 20.4 2805 3405 46.0 52.0 39.0 57.0 6.0 2800 39.0 5280 1.2 2019 2104 2276 2288 20.4 2805 3405 55.0 1.2 2019 2104 2276 2288 20.4 2805 3405 55.0 1.2 2019 2104 2276 2288 20.4 2805 3405 55.0 1.2 2019 2104 2276 2288 20.4 2805 3405 55.0 1.2 2019 2104 2276 2288 20.4 2805 3405 55.0 1.2 2019 2104 2276 2288 20.4 2805 3405 55.0
Rate 10.0 2260 4535 10.0 2260 4535 41.0 46.0 40.6 47.6 40.6 47.6 40.6 47.6 40.6 47.6 40.6 47.6 40.6 47.6 40.0
Grouth Rate 10.0 2260 4535 10.0 2260 4535 48.9 3085 4705 48.9 3085 4705 48.9 37.9 3170 5206 47.5 3170 5206 47.6 48.0 27.9 3121 5192 27.9 3121 27.
Growth Rate 9/day 10.0 2260 48.9 48.9 28.3 227.9 3121 22.5 40.0 48.0 48.1 22.5 1986 3206 46.1 22.5 1986 3206 47.6 20.4 20.4 20.4 39.6 47.6 34.7 38.2 39.9 48.0 48.1 22.5 39.9 48.0 48.1 22.5 39.9 48.0 48.1 22.5 39.9 48.0 48.1 22.5 39.6 40.6 47.6 20.4 38.6 39.7 38.7 38.7 38.7 38.7
4535 50.0 4795 47.6 47.9 4334 48.0 5192 48.0 5192 48.0 5192 48.0 5192 48.0 5420 47.4 48.0 5420 47.6

* Four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

Appendix 1

.•		Vgt_57	4300	4800 57.0	5787 56.4	•	6900 57.3	5600 56.9		6755. 57.6	7150 57.6		6090 57.72	5930 57.7	6256 57.3
		Mgr_48	3900	3750 48.0	4170		5265 48.1	4205		5115 48.0	5100 48.6		4420	4375	
		Ngt 40	2880		2370 39.6	3291 39.6	3335	3310		3280 39.9	3050		3004	2850	3873 42.9
	Growth	g/day	29.3	55.6	30.8	36.7	36.8	42.8	17.71	36.9	43.2	39.6	36.7	35.8	39.2
		Ngt9										1938 33.6			٠
		WgtB										1882			
alyses		Ngt7										1858 33.3			
tical An		Vgt6		2170 35.9								1811 33.1			
e Statis		Ngt5	2140 35.9	2020 34.7	2330					2570 36.0	3050	1778 33.0		•	
ed in th	٠	Ngt4	1960 35.0	1760	1843 36.0	2240 34.0	2260 36.0			2400 35.4	3050	1732	3004	2850 39.3	
includ		Wgt3	1730 34.1	1550 32.7	1616	1980 33.1	1915 34.7	2160 36.3		1990 34.0	2260 38.1	1699	3004	2850 39.3	2500 37.0
f Weights	•	Wgt2	1570 33.1	1370	1446 34.0	1770 32.1	1655	1908	1429	1740	2040	1675 32.6	2045	1923 35.7	1740 34.3
listing of Weights Included in the Statistical Analyses		Wgt1	1380 32.1	1320	1380 33.0	1490	1490	1604 34.4	1305	1555 32.0	1728 36.1	1649 32.4	1780 34.4	1651	1485
-		Variable	s pca)	Veight (9) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	. Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (9) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g)	ueight (g) Age (weeks pca)	Weight (g)
		Subject		9704-0306	9705-0303	9705-0305	9060-9076	9706-0306	9707-0001	9207-0304	9707-0306	9707-0307*	9707-1308	9707-2308	9708-0302
			Gender keyimeri Hale DHA	DIIA	DIIA	DIIA	DIIA	. Alla	DHA	DHA	DHA	DHA	DIIA	DIIA	DHA
		-	Gender Hale	Hal e	Hale	Hale	Male	Hale	Hale	Male	Hale	Hale	Male	Male	Hale

* four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

opendix 1

Listing of Weights Included in the Statistical Analyses

									•					
	Wgt_57	6750 56.4		7300	. \$860 \$7.6			6646 56.6	7937 57.3	4993	5050 57.6	7380	6600 56.7	
	Wgt_48	\$080 47.4		5200 48.1	4680	5500	5840 50.6	\$525 47.6	47.6 47.6	3404 48.0	4256	5540	5055 46.7	5200 48.4
	Mgt_40	3150 39.4		3160	3040 39.6	3100	3628 38.1	2440 37.4	3553	2355	2610	3255 39.7	3240 39.7	3960 42.3
Rafe	g/day	4.4	7.1	30.5	33.9	31.1	32.2	20.9	32.0	29.8	17.2	40.7	48.9	41.4
	Vgt9													
	Wgt8													
	Ngt7													3228 37.7
	Ngt6													3072 37.3
,	Hgt5	2800 36.7		2550 37.6			2440 36.4		•			2735 37.9		2756 36.3
	Ngré	2400 35.4		2160 36.0	1945 34.5	2300	2375 36.0		2120	2355	1490	2570 36.9	2835 37.7	2460 35.3
	Vgt3	34.4		1985 35.0	33.5	2100	2160 35.0	1550 33.6	1870	1950	1290 34.0	2235	2045 35.6	2245 34.4
	Wgt2	1740 33.4	1520	1800 34.0	1435 32.5	1810	1880 34.0	1340	1690 32.4	1689 37.1	1134 33.0	1880 34.7	1686 34.6	2037
	Wgt1	1490	1470	1545	1240	1700	1530	1120	31.1	1499 36.1	1056	1635 33.9	1442	1587
	Variable	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)
	Subject	9709-0301	9709-0304	9712-0304	9712-0306	9743-0303	9743-0304	9698-0305	9050-9696	7020-6696	50£0-6696	DIIA+ARA 9700-0302	9701-0302	9701-0306
	Regimen	ОНА	DIIA	DIIA	DIIA	DIIA	DIIA	DHA+ARA	DIIA+ARA	DHA+ARA	· DIIA+ARA	DIIA+ARA	DIIA+ARA	DIIA+ARA
	Gender	Male	Hale	.Hale	Male	Male	Male	Hale	Male	Kale	на! е	Hale	Male	Male

* four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

Appendix 1

Listing of Weights included in the Statistical Analyses

	49t_48 49t_57	5930 7475 48.6 57.4	5250 47.6	5160 6520 48.0 56.4	6020 6720 47.4 56.6		5460 7050 47.7 56.7			5447 6809 47.9 56.9		4820 6225 48.1 58.1		5225
-	49 ¥9 ¥9	3445 5 40.6	3780 5 40.6 4	3500 5 40.0 41	4350 60 40.4 47	3170 43 40.0 47	3220 54 39.9 47	2570 65 40.0 48	2979 44 40.1 48	3631 54 39.9 47	3007 55 39.9 48	2695 48 39.9 48	3585 5955 40.4 49.1	3460 5255
Growth	-	42.5	36.0	2.05		34.1	35.1	22.2	. 27.0	32.7	36.4	31.4	40.0	40.3
	Wgt9	•												
	WgrB	-												
	Wgt7													
	Wgt6					2590 36.9		1840 36.9						
	HgtS			•	2415 33.4	2390	2050 34.4	1680 36.0		2300				
	H914	2932 38.4	2660 36.0		2055	2115 35.0	1740 33.4	1520 34.9	1870 35.7	2020 34.4	2240 37.4	1930 36.6	2270 35.1	
	Hgt3	1919	2160	2660	1745	1830	1490 32.4	1370 34.0	1620 34.7	1700	1810 36.1	1660	1825 33.9	2150
	Hgt2	1710	1865 33.0	1905 33.9	1460 30.4	1635 33.0	1270	1230 33.0	1440 33.7	1490 32.4	1650 35.4	1455	1585 33.0	1910
	Wgt1	1397	1670	1650	1255	1440	1110	1080	1300	1320 31.4	1480 34.4	1330	1355	1620
	Variable	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Welght (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g)			
	Subject	9701-0307	9702-0301	9702-0303	9703-0301	9703-0305	9704-0301	9704-0302	9705-0301	9705-0306	9705-0307	9706-0305	9706-0307	9706-0309
. ,	Regimen	DHA+ARA	DIIA+ARA	DIIA+ARA	DHA+ARA	DIIA+ARA	DIIA+ARA	DHA+ARA	DHA+ARA	DHA+ARA	DIIA+ARA	DHA+ARA	DIIA+ARA	DIIA+ARA
	Gender	нае	Hale	Male	Male	Hale	Hale.	Hale	Hale	Hale	Ha≀e	Hale	Hale	- A

* Four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

Listing of Weights included in the Statistical Analyses

Gender	Regimen	Subject	Variable	Vgt1	Wgt2	Ngt3	Ngté	Wgt5	Ngt6	N9t7	Wgt8	Wgt9	Growth Rate g/day	05 16M	95 16M	Var 57
Male	DHA+ARA	9707-0301	Veight (g) Age (weeks pca)	1553	1980	2280 35.3	2720 36.6				•		41.5	3395	4950	6285
Hale	DIIA+ARA	9707-0305	Veight (g) Age (weeks pca)	1755 33.9	1990	35.7	2505	2770 37.7					37.4			
. Hale	DIIA+ARA	9707-0310	Veight (g) Age (weeks pca)	1620 32.7	1828 33.7	2140 34.7	3195	•					44.8	3585 39.7	5170 47.9	6725
Hale.	DIIA+ARA	DIIA+ARA 9708-0301	Weight (g) Age (weeks pca)	1640	1880 33.7	2200 34.7	2420 35.7	•					38.0	3730	4835	6185
Hale	DHA+ARA	9708-0304	Weight (g) Age (weeks pca)	1680	2180 35.9			:					55.6			
Hale	DIIA+ARA	DIIA+ARA 9709-0303	Weight (g) Age (weeks pca)	1470	1810								48.6			
Hale	DIIA+ARA	9709-0305		1410 34.4	1655	1900 36.4	2160						35.6	2630 39.7	4570	5520 57.1
Male	DHA+ARA	9712-0303	Weight (g) Age (weeks pca)	1180	1210 32.3	1450 33.4	1590 34.4						6°02	2520 40.4	3500	5010 56.4
Hale	DHA+ARA	9712-0305	Weight (g) Age (weeks pca)	1325	1505 32.5	1785	2010 34.5	2300					34.1	3030	4350	5510 57.6
Hale	DIIA+ARA	9723-0301	Weight (g) Age (weeks pca)	1630 33.9	1728 34.9	1961 35.9	2214 36.9						28.4	3104	•	5986 58.9
Hale	£	1090-8696									•			3518 40.0	5497 48.3	6582 56.9
Hale	¥	9698-0602		·										3177	\$220 48.1	6355 57.0
Male	¥	8090-8696		· .										3858 40.0	5447 48.0	6454 57.0

Four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

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Appendix 1 Listing of Weights Included in the Statistical Analyses

		₽.			-							•	Growth			
Gender	Regimen	Subject	Variable	Vgc1	Hgt2	Wgt3	H9t4	HgtS	Ngt6	Wgt7	Wg t 8	Ngto.	kate 9/day	Ngr_40	Hgt 48	Har 57
Male	£	7090-8696		: .										4355	5005	6383
- E	=	50901-8696				•								0.04	48.0	57.0
-								,						3433	4979	6426 57.1
9	<u> </u>	1000-6606												3915 40.0	6639 48.3	7775 57.72
Hale		7000-6696												3802 40.0	5787	7178
a e .		1000-1074												3317 40.0	5555	7070 56.4
Hale	ž i	7000-1076						•						3487 40.0	5833 47.3	8070 58.3
Hale e	≚ :	5000-1076												3232 40.0		5855 56.4
Ha!e	<u> </u>	700-10/6												3600		6285 56.9
Hale	Ī	5090-10/6		-									٠	3402 40.0	•	7210 57.6
πa e		9701-0000					٠							3090		5445 56.7
a .		1000-2016						•						3480 40.0		6530 56.6
Hare -	<u>.</u>	2000-2016	•	٠.	•			·		•				3165 40.0	5060 48.3	6660 57.1
Hale	Ē	2000-5078												2670 40.0		7220 57.1

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Listing of Heights Included in the Statistical Analyses Appendix 1

													Rate			
Japan	8 e a i men	Subject	Variable	Ngt1	Wġt 2	·Vgt3	Ngt4	Vgt5	Wgt6	Ngt7	WgtB	Ngt9	g/day		87 JGH	49t_57
															7 L7 07L9	8330
		er condu-														7.
Male	¥	9703-0504		 1:										3435 40.0	6000 48.1	7930 57.1
Male	H.	9704-0502												3285	5220 48.1	6560 56.6
Hale	. · =	9704-0503												3400	5200 48.7	6725 56.9
Male	H	9705-0601		·										3200	5617 48.3	6752 57.3
наве	Ħ	9705-0602												3860	6227 48.0	
Hale	¥	9706-0601												3152 40.0	\$105 49.0	6545 .57.0
Hale	蓋	9706-0602												3557	5175 47.4	7315 57.72
Hale		9706-0603						-						3192	5070 47.9	6970 56.7
Hale	폴	7090-9026												3461	4225	5525 57.1
Male	¥	9706-0605												3870 40.0	6220	7660 56.4
Hale .	N	9090-9026	2	•							-		-	4315	5975	6720 56.6
Male	¥.	9707-0601	_											3263 (0.0)	4730	5825 57.0

Listing of Weights Included in the Statistical Analyses

Gender	Regimen	Subject	Variable		Wgt1	Hgt2	Hgt3	.536H	VgtS	Ngt6	Ngt7	WgtB	Ngt9	g/day	Ngt_40	Ngt 48	Hgt 57	
Hale	Ħ	9707-0602										·			3206 40.0	4515	- 6220 57.7	
Hale	¥	9707-0603			.,										4256	6930 48.0	8810 57.0	
Ha∫e	Ŧ	9090-2026													34.19	5460 48.0	6130 56.7	
Hale	Ξ .	9707-0605		٠.					÷						3433			
Hale	₹ .	9707-0606		• • .											3603	5825 48.4		
. на е	¥	7090-1046				•									3569 40.0	5410 47.9	6870 56.9	
Hale	≡	9707-0608													3348	5135	6370 57.0	
Male	Ξ ·	6090-2026											٠		3348			
Hale	폴	9708-0601													3064	5220 47.6	6595 56.4	
Male	豎	9708-0602			•										4085 40.0			
Hale e	Ħ	9708-0603													3319 40.0	5135 48.4	6327 57.1	
Hale	Ŧ.	9708-0604								•					3291			
Hale	포	9708-0605													3796			

• four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

ppendix 1

Listing of Weights Included in the Statistical Analyses

Ngr_57	5405 57.1	5180 56.7			5220 56.9	5816 57.0	5200 55.6	6280 56.4	5815	5505 57.4		6900 56.7	97.75 57.6
49 £ 48	4645 48.4	4043 47.9			4369	4596 48.0	4165 48.6	\$140 48.4	4425	4420	47.6	5160 47.7	4820
Wgt_40	4050 40.0	3333	3400		2610 39.7	2780	2675 40.6	3175 39.7	2980	2870 39.7	3380	39.9	3060 39.9
Growth Rate g/day				5.6	24.1	37.3	29.1	28.3	41.1	36.6	7.62	31.6	42.2
49£9							,						
Идгв				1070 32.1									
Ngt7				1080 32.0							٠		
Ngté				1060 31.9							2390		
V9t5				1080	2145		2292 38.6	1976 34.7	2406 37.9	2044	1995 36.0		
Ngt¢				1080 31.6	2000 35.7	24 <i>97</i> 38.0	1975	1745	2198 37.3	1756	1750 35.1	2530 36.0	2645 37.0
H9t3	•			1070	1862 34.7	1860 36.0	1903 36.6	1555	1898 36.4	1492 32.4	1570 34.1	1840 33.1	2410
Wgt2				1050 31.3	1672 33.7	1629 35.0	1633 35.6	1366	1569	1254	1371 32.7	1555	2065
Wgt 1		٠,		1020 31.1	1464 32.7	1473 34 0	1480 34.6	1174	1391	1050 30.6	1222 31.7	1454 31.0	1775 34.0
Variable				Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca).	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)
Subject	9090-8026	7090-0076	9709-0505	*E000-8696	1000-6696	. £000-6696	9701-0003	9701-0005	9701-0008	9701-0011	2000-2026	900-2026	Control 9702-0010
Reginen	¥	¥	¥	Control	Control	Control							
Gender	Hale	Male	. Hale	Female	Female	Female	Female	Femate	female	f ema l e	Female	f ета l е	Female

Female

. Female

Female

Female

Female

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4330

4085

4gt_57

Gender Female female

female

female

Female

		40 Agr_40	3210 39.6	2610 37.3	3360	2722 39.7		2740 40.0	3640	3655 40.0	2680	3320	3110 40.1	3430	3330
	Growth	g/day	26.4	29.5	48.3	28.3	37.9	31.7	31.6	56.0	31.1	32.6	30.2	41.2	39.9
		Vgt9													
		Ng t 8													
Analyses		Ngt7													
Istical		Wgt6	2130 34.3										2765		
the Stat		WgtS	1825 33.4	2220 35.3	2685 36.6								2325 36.4		
Listing of Weights Included in the Statistical Analyses		Hgt4	1570	1900 33.9	2445 36.0	1660 34.0	2330	2150 36.0			1810 34.6	· · · · · · · · · · · · · · · · · · ·	2010 35.3		
hts Incl		Vgt3	1390	1765 33.3	2095 35.0	1490	1965	1805	1960 34.3		1585 33.6	1935	1655 33.6	3430	3330
of Weig		Wgt 2	1250 30.4	1590	1715 34.0	1290 32.3	1673 36.3	1610 33.7	1620	2185 35.0	1270 32.4	1765 33.1	1505	3430	3330
Listing		Wgt1	1170 29.1	1420 31.4	1495	1120	1515 35.1	1485 33.0	1525	1905	31.6	1510	1465 32.0	1866 34.6	1815
		Variable	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Height (9) Age (Heeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)			
		Subject	9703-0002	9703-0005	9703-0008	9000-5026	9706-0003	9706-0005	9706-0009	9706-0010	9706-0013	9706-0016	9707-0003	9000-2026	9707-1006
		Regimen	Control	Control	Control	Control									

6535 56.7

5390 48.4 3800 48.4

5495 47.6 529*7* 56.6

4835

4995

female

Female

female

Appendix 1

Listing of Weights Included in the Statistical Analyses

	Wgt_57			5361 57.1		5900 56.7	5880 57.1	6230 57.0	5160 57.4	5192	6291 57.0		5121 57.0	5699 56.4
	85 16H	4734		4110		4700	4450 47.6	4560 48.0	7.87 78.4	4145	48.0		3979 48.0	5185 47.4
	Ngt 40	. 2910 40.6		2582 39.3		2975 39.6	2930 40.3	3030	•	3170 39.7	3787		2891 40.0	3135
Growth Rate	g/day	27.2	4.3	33.1	30.0	32.3	25.6	28.4	24.0	42.7	34.7	28.7	55.9	29.7
	Mar9													\
	Мдгв													
	Vgt7													
	Wgt6													
	Hgt5					2200 36.3	1695 33.1			2625 36.6			2020 34.9	
	V9t4	2050 36.9		2180 36.3		1945 35.6	1490 32.1	1790	1890 35.1	2320	2140 33.1	1720 33.7	1630 33,1	
	Ngt3	1850 35.4		1860	24.00	1665 34.6	1290	1585 33.9	1740	2075 34.6	1890 32.1	1420 32.7	1520 32.4	2450
	Ng t 2	1600 34.4	970.0 31.0	1605	33.7	1425	1145	1358	1520	1740 33.6	1650 31.1	1240	1310	2110 35.7
	Wgt1	1410 33.4	940.0 30.0	1380			972.0 29.1	1203	1300					1790 34.4
.:	Variable	Weight (g) Age (weeks pca)	Veight (g) Age (veeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (9) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)				
	Subject	9708-0001	9708-0003	9708-0008	9709-0002	9709-0005	Control 9712-0005	9000-2126	9743-0003	9746-0001	9000-8696	9000-8696	6000-8696	9698-0307
	Regimen	Control	DIIA	DIIA	DHA	DHA								
	Gender	female	female	female Control	Fenale	Feinale	Female							

* Four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

Appendix 1

Listing of Weights Included in the Statistical Analyses

													Grouth			
	٠.		•									•				
	,	1. 1. 1.	Variable	Vgt 1	Hgt2	Wgt3	Hgt¢	Ngt5 1	Hgt6	Vgt7	₩gt8	4919		Ngt_40	Ngt_48	Wgt_57
Gender Female	Regimen		Weight (9)	1313	1477	1669 34.9	1929 35.9	2380 36.9					36.9	39.7	7.72	7093 56.7
Fenale	OIIA	9700-0001	Meight (9)	1580	1820	2050	2295	2500 36.3					34.5	3210 40.1	5110 48.1	6300 57.1
Female	·DIIA	1000-1026	Age (Necks post) Height (9) Age (Heeks post)	1300	1356 34.0	1586 35.0	1924 36.0	2125 · 36.6					34.2	2910 39.6	4325 48.0	\$625 \$7.0
female	рна	9701-0004	Weight (9) Age (weeks pca)	1108	1261	1441	1671	1897 34.7					28.4	3020 39.7	4855	56.4
Female	DHA	9701-0012	Weight (g) Age (weeks pca)	1674	1928 35.9	2151 36.9	2311 37.6	2685 39.6	2685 39.6				30.1	2685 39.6		
Female	DIIA	9701-0014	veight (g) Age (weeks pca)	1422	1631 34.9	1858 35.9	2455						37.2	39.9	4605	5140
Female	DHA	9702-0001	Weight (g) Age (weeks pca)	1780 31.6	2115	2390	3000						35.8	3850	\$610 49.6	6600 57.0
Female	DIIA	9702-0006		1850	2005	2650 39.6	2650 39.6						27.3	2650 39. 6	4450 48.4	6020 56.4
Female	e DHA	9702-0007	Height (g) Age (weeks pca)		1459 32.1	1780 33.6	1965	2035					29.6			
female	e DHA	9702-0008	Weight (9) Age (weeks pca)	1605	1930	3540 39.6	39.6						51.3	39.6	2850	7820 57.1
Female	e DIIA	9703-0003	Weight (g) Age (weeks	1255	1355	1535	1845 37.1	2150 38.1					34.8	39.4	4130	5010
Female	Le DNA	9703-0004		1170	1340	1550	1795	2225 37.0					33.9	39.4	4610	57.1
Female	le DHA	9703-0009		1570	1830	2095 35.1	2395	2655 37.9					34.6	5160 40.4	7.87	58.0

Appendix 1

Listing of Weights included in the Statistical Analyses

	49t_48 4gt_57	5830 8630 48.0 57.0	4860 6100 48.0 57.0	4795 5986 48.1 57.1	(145 5320 (8.1 57.3		4790 48.4	5600 7675 49.4 58.0	4595 5765 48.0 57.0	•		4620 6530 48.1 57.0		4080 5420
	Ngt_40 Wg	3100 5	3360 4 39.6			2120 39.9	3530 4 40.1 4	3295	3045 4			3010 4 40.1 4	3500	2580 4
Growth Rate		30.5	30.0	31.9	31.7	23.0	32.5	26.2	38.1	42.2	38.1	39.5	33.8	30.5
	Wgt9													
	NgtB													
	Ngt7													
	Wgt6												2520 35.0	
	Vgt5		1890 33.7	2098 36.7			•	1804			2485 36.6		2250 34.0	
•	Ng t 4		1700 32.7	1880 35.7	1930 36.0	1485 36.4	34.6	1560 34.4			2280 35.6	2380	1970 33.0	2155
	Ngt3	1740 35.0	1490 31.7	1590	1630	1345 35.7	2130	1395	2850 38.7	3440	1955	2110 35.7	1755 32.0	2015
	Hgt2	1670 34.6	1310 30.9	1370 33.6	1405 33.7	1188 34.6	1830 32.4	1170 32.6	1771 35.0	3440	1665 33.6	1775 34.7	1490	1725
	.Wgt1	1440	1050	1220 32.7	1270	990.0	1610 31.6		1635			1485	1250	157.0
	Variable	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	. Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	
factoria i e	Subject	5000-5026	\$000-5026	1000-5026	9000-9026	9706-0008	9706-0012	9706-0014	9707-0004	9707-0308	9708-8076	9000-8026	1000-6026	। इन हैं
	Regimen	DHA	риа	DIIA	DIIA	DIIA	DIIA	DHA	рна	DIIA	DIKA	DIRA	DHA	
	Gender	Female	Female	Female	Female	Female	female	Female	Female	Female	Female	Female	Femate	

* Four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

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Listing of Weights Included in the Statistical Analyses

	Vgt_57	5250 57.1		5340 56.9	5,72	5160 57.4	6582 56.7		6979 57.3	8341 57.0	6420 57.1	6525 56.4	6270 57.3	6955 56.9
	Wgt_48	3980 48.1		4250	4140	4540	5348		5107 48.3	6752	4930	5115	5045	4935
	Ngt_40	2940 40.1		2425 39.7			3530	3241 40.7	3177	4029	3340 40.3	2930 39.4	3600	2680 39.9
Growth	g/day	54.9	56.4	27.3	33.5	29.7	37.1	31.8	28.9	35.1	31.9	37.8	38.3	29.8
	Wgt9							•						
	Wgt8													
	Hgt7													
	Wgt6										2480 35.6			
	Wgt5	1685 34.0			1930				1788 35.0	2330	2220 34.1	2420	2728 36.1	2227 37.7
	Hare	1470		1650 35.7	1800 35.7	1975	2360 34.9	2260 35.7	1536 34.0	34.9	2035	2210 36.4	2456 35.3	1982 36.7
	.Wgt3	1270 32.0	1430 34.7	1440 34.7	1470 34.4	1845 34.1	33.6	2130 34.6	1283 33.0	1688 33.9	1885 32.3	1887 35.4	2113	1590
	Wgt2	1120 31.0	1230 33.7	1230 33.7	1170 33.1	1570	1690	1870	1122 32.0	1542 32.9	1525	1609 34.4	1859	1427 34.9
	. Wgt1	987.0 30.0	1060	1082 32.7	1000	1380	1550 31.6	1580 32.6	985.0 31.0	1330	1315	1398		1469
	Variable	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks.pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)			
	Sub Ject	9712-0001	9712-0002	9712-0007	1000-5726	9743-0002	9698-0001	9698-0002	5000-6696	5000-6696	9700-0005	9701-0002	9701-0006	9701-0007
	Regimen	DHA	DIIA	DIIA	DIIA	DIIA	DIIA+ARA	DIIA+ARA	DHA+ARA	DHA+ARA	DHA+ARA	DHA+ARA	DHA+ARA	DIIA+ARA
•	Gender	Female	Female	female.	female	Female	Female	Female	female	. Female	Female	Female	female	female
·				•							•			

* Four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

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Listing of Height's Included in the Statistical Analyses

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Wgt_57		5550 57.4	7500 56.9	5340 56.4	6410 57.6	5420 57.3	6650 56.7	5850 56.7	6800 57.3	6640 57.0	6894 56.9	5050 57.0	7655 56.7
Wgt_48	5545 48.4	4545	6220 48.4	4300	4680	4250	5400 48.1	4190	5150 48.0	5400 48.0	5107 48.4	4000	6550 48.6
05 ⁻ 16H	3500 41.1		4190	3025	2905 39.9	3030	3600	2850	3110	0.07 0002	3376 39.9	2600 40.4	4100 40.1
Growth Rate 9/day	34.6	35.6	39.9	59.9	6.02	28.9	49.1	27.4	26.7	30.0	8.67	22.1	34.5
Vgt9										•		1380 33.4	
Wgt8												1350 33.3	
Hgt7	,								2070 34.9			1265 33.0	
. 916M	2759 37.7							2240 36.6	1780 33.9			1310 32.7	
Wgt5	2433 36.1		2400 34.1	2710 38.0	2655 37.3	1955 35.3		2030 35.7	1570 32.9	·		1310 32.4	
7161	2234 35.3	•	2155	2525 37.0	2595 37.0	1680 34.3	2880 37.0	1880 35.0	1370		2920 37.7	1280 32.1	2060
Ngt3	1978 34.4		1820 32.1	2300 36.0	2230 36.0	1450 33.1	2560 35.9	1620	1200		36.6	1185	1685 33.7
Vg t 2	1703	2019	1488	2060 35.0	2000 35.0	1255 32,1	2200 35.0	1495	1090 30.0	1840 33.4	2260 35.7	1120	1515 32.9
Wgt1	1488 32.3	1841 33.0	1293 30.1	1895	1725 34.0	31.3	1865	1390	960.0	1690	1760 34.4	1075 31.1	1290 31.7
Variable	Veight (g) Age (weeks pca)	Weight (g) Age (нееks pca)	· Veight (g) Age (weeks pca)	Veight (9) Age (weeks pca)	Veight (g) Age (weeks pca)	Veight (g) Age (weeks pca)	Ueight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Weight (g) Age (weeks pca)	Veight (g) Age (weeks pca)			
Subject	9701-0010	9701-0013	9702-0003	9702-0005	6000-2026	9703-0001	9000-5026	9703-0007	2000-5026	DIIA+ARA .9704-0003	9705-0003	*5000-5026	9706-0001
Regimen	DHA+ARA	DHA+ARA	DIIA+ARA	DIIA+ARA	DHA+ARA	DHA+ARA	DIIA+ARA	DIIA+ARA	DHA+ARA	DIIA+ARA	DIIA.+ARA	DIIA+ARA	DHA+ARA
Gender	f emal e	Female	female:	Female	female	Female	f emal e	Feinale	Female	Female	f emal e	Female	Female

		··· - <u>-</u> -				νbί	Appendix 1										
				Listing of Weights Included in the Statistical Analyses	of Veigh	ts Inclu	ded in tl	he Stati	stical A	Analyses					-		
Gender	Regimen	Subject	Variable	Wgt 1	Vgt2	Wgt3	Hgt4	Hgt5	Ngté	Hgt7	WgtB	Wgt9	Growth Rate g/day	Ngt_40	49t_48	Wgt_57	
Female	DHA+ARA.	9706-0002	Veight (g) Age (weeks pca)	1395	1710 33.0	1884 33.9	2275 35.4						34.8	2845	48.9	5550 57.3	
f emale	DIIA+ARA	5000-9026	Veight (g) Age (weeks pca)	1550 36.7	1705 37.6	2050 38.7							36.1	2645	4225	6935 58.0	
Female	DHA+ARA	7000-0076	Weight (g) Age (weeks pca)	1235	1490	1820 35.7	1930 36.4						34.3	2505			
Female	DIIA+ARA	9706-0011	Veight (g) Age (weeks pca)	1900 34.3	2105								41.0	3430	5175	6140 56.7	
female	DIIA+ARA	9706-0015	Weight (g) Age (weeks pca)	1670	1975 35.6	2210 36.4							41.6	3005	4465 48.4	5810 57.6	
Female	DHA+ARA	DIIA+ARA 9706-0017	Weight (g) Age (weeks pca)	1465	1700 33.4	1895 34.3	2170 35.3						33.4				
f emale	DHA+ARA	9707-0002	Weight (g) Age (weeks pca)	1775 34.3	2240 36.0	2385 36.9	2610 37.9						33.2				
female	DIIA+ARA	9708-0002	Weight (g) Age (weeks pca)	1535 33.0	1700	1980 35.0	2200 36.0					•	. 32.5	2724 38.1	9.72 47.6	6315 55.4	
Female	DIIA+ARA	9708-0005	Weight (g) Age (weeks pca)	1125	1345	1610 34.4	1980 35.4						7.07	3121 39.4	5855	7875 57.4	
female	DIIA+ARA	7000-8079	Weight (g) Age (weeks pca)	1200	1440	1680 33.3	1975 34.3						36.6				
Female	DHA+ARA	600-6026	Weight (g) Age (weeks pca)	1350	1560 33.3	1885 34.6	2250 35.6	2475 36.3			٠		37.0	3295 39.7	5250 48.4	56.85	
Female	DHA+ARA	9712-0003	Veight (g) Age (неeks pca)	1283 32.0	1410 33.0	1590 34.0	1850 35.0	2010 36.0					27.1	2580	4130	5640 57.5	
female	DHA+ARA	9712-0004	Veight (g) Age (weeks pca)	1575 33.0	1760 34.0	1890	2080	2530 37.6	•				29.7	3220 40.3	4920	6600 57.1	
			•														

* Four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

Appendix 1

Listing of Weights Included in the Statistical Analyses

•	Wgt_57	5760	5362			5319	6,72	\$653 \$7.0	5731 57.0	5986 57.0	5674 56.7	6355 57.0	57.6	8450 57.7
	Wgt_48	4470	4010	4880	5972	4213	5234 48.7	4638 48.3	4766	4823	4482	4738	5617 48.4	5630 47.7
	Ngt_40	2960 40.1	2680 39.9	3546 40.0	3518 40.0	3390	3383	3646 40.0	2582 40.0	4284 40.0	3716 40.0	3660 40.0	3433	3884 40.0
Growth Rate	g/day	37.2	30.1											
	4364 4									•	٠			
8	2										,			
Vor 7														
Vaté	•													
Wats			2110 36.7											
Wgt4	, ;	37.4	1814 35.7											
Hgt3	•	35.8	1597 34.7		•									
Ngt2	007	35.0	1429						•	•				
Wgt1	. 60	34.0	1249 32.7					•						
:		s pca)	(eod s			•							. *.	
Variable	11. 20.0	метупт (g) Age-(weeks pca)	Weight (g) Age (weeks pca)			,								
Subject	פטטט ריינט	9000-7176	9746-0002	9698-0501	9698-0502	9698-0503	9698-0504	\$050-8696	9699-0601	7090 - 8696	5000 - 6606	7090-6696	9699-0605	9701-0501
Gender Regimen	4	OHA+AKA	DHA+ARA	¥	¥	Ξ	≚	¥	¥	Œ.	≝	₹	¥.	Ŧ
Gender	· .•	remale	f emale	Female	f ema l e	female	Female	female	f emale	f ema l e	Female	female	f emal e	female

* Four subjects had more the 9 weights used in growth rate calculation. A complete listing appears on the last page.

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Listing of Weights Included in the Statistical Analyses

•		V.				,	ı							Growth			
Gender	Gender Regimen	Subject	Variable		Wgt1	Ng t 2	N9t3	Ngr4	Vgt5	Ngt6.	Vgt7	WgtB	Wgr9	g/day	05 16N	Wgt_48	Wg (
Female	¥	9701-,0502													3858	5420 48.6	670 57.
Female	¥	9701-0503												,	3430 40.0	4265	508! 57.4
female	Ξ	9701-0504							,						3317	5020	623(57.
f emal e	¥	9702-0501													3302	5540	6630 56.7
Female	Ξ	9702-0502	•	٠,		٠.									2658 40.0	5310 47.4	6800 57.1
female	Ŧ	9702-0503													2895	3430	4530
f enial e	Ŧ	9702-0504													3401	5390	6270 57.4
Female	¥	9702-0505				•							•	•	3141	4210	5320 57.0
f emale	Ħ	9702-10506													3762	6040 48.9	7600 57.7
Female	H	9702-0507					٠							•	2718 40.0	4050	4940 57.4
Fenale	¥	9702-0508													2927 40.0	4540	5860 57.0
Feniale	W .	9703-0501										-			4085 40.0	5260 48.1	6360 57.1
Female	Ŧ	9703-0505										•	-		3390	5760	7670

Listing of Weights Included in the Statistical Analyses

	Wgt_S	7490	6550 56.3	5880 57.4	5702 57.1	7348 57.3	6645 58.1	5525 57.6	6770 56.6	7080 57.1	7675 56.9		6890 57.6	5950 57.4
	Mgt_48	6170	5090 48.0	4700	4500	6327 48.3	5000	4315	5515 47.9	5500	5785 47.9		5440 48.1	
	Wgt_40		3085										3688 40.0	
Growth Rate	g/day							٠					•	
٠.	Ng t 9									•				
9	Mgt8													
7	/ 16M											,	٠	
7407	0							•.						
, you	2	•												
7100	, ,					٠								
Vor3					ć									
Wat 2							•						٠	
Wat 1											•			
	₹.												,	
Variable														
Subject		ar B	9703-0507	9/04-0501	1060-6078	2050-5078	9/06-0501	7050-90/6	1050-7076	7050-7076	5050-7076	9707-0505	9708-0501	2050-90/6
Gender. Regimen	¥		Ξ :	ž i		٠.	·		X	E :	Ē	₹		Ĕ
ander.	a lema i											Female		Female F
ٽ	ŭ		ب ن	Ξ ,	Ξ.	Ξ,	Ξ ,	Ξ ,	I (Ξ,	Ĭ	7	<u>.</u>	<u>~</u>

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			40.000	· .		Ç	<u> </u>	7			:			Growth			
ender	Regimen	Subject Subject	Variable		- 164	216H	C16#	536A	CJ 6M	Hato	Mgt7	Mg t 8	H919	g/day	Mgt_40	Hgt_48	Hgt_57
етаве	¥	9708-0503	•												2977	5165	7040
emale	¥	9708-0504					-								3864	5660 48.4	6705 57.4
emale	¥.	9708-0505		٠.											3831 40.0	5800	7435
emale	¥	9709-0501				·	-								3550 40.0	•	
emale	¥.	9709-0502	`	. ',		,									3715 40.0	5205 48.0	6100 56.9
enale	· E	9709-0503					•								3195 40.0		•
female	王 ·	9709-0504													3190 40.0	4590	
emale	至	9709-0506											•		3505 40.0	4500 48.0	5910 57.1

Appendix 1
Listing of Weights Included in the Statistical Analyses

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Listing of Weights Included in the Statistical Analyses

Rate 9/day	26.1	39.6	5.6	22.1
				1670
Wgtio Wgtii Wgti2 Wgti3 Wgti4 Wgti5 Wgti6 Wgti7 Wgti8				1680
19t16 1				1640 34.6
/gt15_1				1585 34.4
1911g				1565
lgt13 L				1515
Jgt12 4		2075 34.0		1510 34.0
/gt11 L	1465 33.0	2030		1450 33.9
Jgt 10 1	1448 32.9	1994		1440
Vgt9 1	1433 32.7	1938 33.6		1380 33.4
Wgt8	1402 32.6	1882 33.4	1070 32.1	1350 33.3
Vgt7	1369 32.4	1858 33.3	1080 32.0	1265 33.0
Ngté	1330 32.3	1811	1060 31.9	1310 32.7
Wgt5	1294 32.1	1778 33.0	1080 31.7	1310 32.4
Wgt4	1291 32.0	1732 32.9	1080 31.6	1280 32.1
Vgt3	1245 31.9	1699	1070 31.4	31.7
Wgtl - Wgt2	1221 31.7	1675 32.6	1050 31.3	1120 31.4
Wgt1	1245 31.6	1649 32.4	1020	1075 31.1
	bca)	bca)	pca)	bca)
Gender Regimen SUBJECI Variable	Hale Control 9712-0301 Weight (g) Age (weeks pca)	9707-0307 Weight (g) Age (weeks pca)	female Control 9698-0003 Weight (g) Age (weeks pca)	Female DIIA+ARA 9705-Ö005 Weight (g) Age (weeks pca)
ECT V	0301 6	0307 1	0003 1	1 5000
SUBJ	9712-	-7076	-8696	9705
egimen	ontrol	ΙΙ	ontrol	IIA+ARA
der Ro	ت	e DIIA	nate C	nale Di
Gen	Hal	Hale	Fen	Fen